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**Department of Defense
Fiscal Year (FY) 2011 President's Budget**

February 2010



Chemical and Biological Defense Program

Justification Book

Research, Development, Test & Evaluation, Defense-Wide - 0400

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Chemical and Biological Defense Program • President's Budget FY 2011 • RDT&E Program

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Department of Defense Chemical and Biological Defense Program Overview

Fiscal Year (FY) 2011 President's Budget

The DoD Chemical and Biological Defense Program (CBDP) is a key part of a comprehensive national strategy to counter the threat of chemical and biological weapons as outlined in the National Military Strategy to Combat Weapons of Mass Destruction, February 2006. The military mission is to dissuade, deter, defend, and defeat those who seek to harm the United States, its allies, and its partners through WMD use or threat of use and, if attacked, mitigate the effects and restore deterrence. This mission is in direct support of the three pillars (non-proliferation, counterproliferation, and consequence management) of the National Strategy for Combating WMD. The DoD CBDP provides research, development, and acquisition (RDA) programs primarily to support the counterproliferation and consequence management pillars. In support of counterproliferation, the DoD CBDP provides passive defenses tailored to the unique characteristics of the various chemical and biological weapons, including emerging threats. These capabilities provide U.S. forces the ability to rapidly and effectively mitigate the effects of a CB attack against our deployed forces. In support of consequence management, the DoD CBDP provides capabilities to respond to the effects of WMD use against our forces deployed abroad, and the homeland.

The CBDP funds research to exploit leading edge technologies to ensure that U.S. forces are equipped with world class capabilities to defend against CB threats through the far term. This budget includes support of a comprehensive science and technology base program to assure we have the technologies needed to protect our troops. CBDP Science and Technology (S&T) research provides core capabilities to ensure U.S. technological advantages, including research into advanced chemical and biological detection systems, advanced materials for improved filtration systems and protection systems, advanced decontaminants, investigations into the environmental fate of chemical warfare agents, advanced information technologies, medical biological defense research (including novel biodefense initiatives that focus on interrupting the disease cycle before and after exposure, as well as addressing the bioengineered threat), diagnostics, therapeutics, and vaccines for viral, bacterial, toxin, and novel threat agents), and medical chemical defense (including investigations of low level chemical warfare agent exposures, diagnostics, therapeutics, pretreatments for classical chemical warfare threats and novel threat agents).

Technologies currently in Budget Activity 4 (Advanced Component Development and Prototypes) and Budget Activity 5 (System Development and Demonstration) provide leading edge tools that will enhance CB defense capabilities for U.S. forces in all CB defense missions in the near-term. The response to chemical and biological threats requires tailored approaches that recognize the fundamental differences between chemical and biological weapons (and even the different types of these threats). This budget details the comprehensive array of systems under development essential to support principles of contamination avoidance, protection, and decontamination.

Key systems in Budget Activity 4 and Budget Activity 5 in FY11 include: the Joint Chemical Agent Detector (JCAD) for portable point chemical agent detection, Joint Effects Model (JEM) and Joint Warning and Reporting Network (JWARN) to provide risk management, comprehensive analysis and response capability tools to the Warfighter, Joint Materiel Decontamination System (JMDS) for interior and sensitive equipment decontamination, Human Remains Decontamination System (HRDS), Sensor Suite Integration (SSI) for NBC Reconnaissance Systems (Stryker), Next Generation Chemical Standoff Detection (NGCSD), Chemical, Biological, Radiological, Nuclear (CBRN) Dismounted Reconnaissance Systems (CBRN DRS) providing equipment integrated into a modular, transportable container for enhanced dismounted operations, Common Analytical Laboratory System (CALS), Joint Biological Point Detection System (JBPDS), Joint Biological Stand-off Detection System (JBSDS) Increment 2, Advanced Anticonvulsant System (AAS), Bioscavenger, Improved Nerve Agent Treatment System (INATS), biological defense vaccines (including botulinum vaccine and plague vaccine), Critical Reagents Program (CRP) to support development of reagents for biological detection and diagnostic systems, Joint Bio Tactical Detection System (JBTDS), Joint Expeditionary Collective Protection (JECF), Joint Service Aircrew Mask (JSAM) and Joint Concept Technology Demonstrations (JCTDs).

In FY 2011, the CDBP will start or continue procurement on a variety of CB defense systems intended to provide U.S. forces with the best available equipment to survive, fight, and win in CB contaminated environments. New starts in procurement for FY 2011 include the Non Traditional Agent Detection Program (NTAD) that will enhance the Warfighter's ability to attain situational awareness and respond to unknown and emerging hazards and the HRDS, which will provide the capability for safe intra-theater handling and storage of Contaminated Human Remains resulting from chemical contamination. Programs continuing procurement include the Joint Service Transportable Decontamination System - Small Scale (JSTDS-SS), Joint Service Personnel Decontamination System (JSPDS), the Joint Effects Model (JEM), Joint Service General Purpose Mask (JSGPM), JWARN, Joint Service Protective Clothing (PROT CLTH) technology, CBRN DRS, Joint Bio Point Detection System (JBPDS), biological defense vaccines, CB Protective Shelters (CBPS), Collectively Protected Field Hospitals (CPFH), Joint Biological Agent Identification System (JBAIDS), Collective Protection System Backfit (CPSBKFT), Critical Reagents Program (CRP), and chemical and biological defense equipment for installation force protection.

Overall, the FY 2011 Budget Estimate achieves a structured, executable, and integrated medical and non-medical joint CB Defense Program that balances urgent short-term procurement needs that include securing the homeland from terrorist attack and emerging threats, against the long-term S&T efforts required to mitigate future CB attacks. Two key initiatives continuing in the FY 2011 submit include the Transformational Medical Technologies Initiative (TMTI) and efforts to enhance detection, medical countermeasures, decontamination, and protection capabilities against NTAs. TMTI is a FY06 Quadrennial Defense Review initiative to protect the Warfighter from emerging and genetically engineered biological threats by providing a novel response capability from identification of pathogens to the development of medical countermeasures (MCM). The focus of the FY 2010/11 TMTI profile will shift towards advanced development efforts as selected candidates enter the FDA clinical trials process. NTA enhancements provided in FY 2010 continue into this FY 2011 submit with further efforts directed towards providing near-term capabilities to the Warfighter while at the same time addressing next generation capability needs. NTA capabilities are accomplished through an integrated portfolio across the CDBP focusing on the enabling Science, Technology and Testing and the advanced development of detection, medical countermeasures, decontamination, and individual protection products. In summary, the DoD CDBP remains committed to establishing the optimal balance between the near term requirement to field modernized equipment to the field, and the need to protect and replenish our long term investment in technology.

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Program Element Table of Contents (by Budget Activity then Line Item Number)

Budget Activity 01: Basic Research

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Budget Activity 02: Applied Research

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Budget Activity 03: Advanced Technology Development (ATD)

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Budget Activity 04: Advanced Component Development & Prototypes (ACD&P)

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Budget Activity 05: Development & Demonstration (SDD)

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Budget Activity 06: RDT&E Management Support

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148	06	0605384BP	CHEMICAL/BIOLOGICAL DEFENSE (RDT&E MGT SUPPORT).....	468
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Budget Activity 07: Operational Systems Development

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Program Element Table of Contents (Alphabetically by Program Element Title)

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CHEMICAL/BIOLOGICAL DEFENSE (ATD)	0603384BP	34	03.....	95
CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)	0601384BP	06	01.....	1
CHEMICAL/BIOLOGICAL DEFENSE (OP SYS DEV)	0607384BP	181	07.....	495
CHEMICAL/BIOLOGICAL DEFENSE (RDT&E MGT SUPPORT)	0605384BP	148	06.....	468
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Chemical and Biological Defense Program
 FY 2011 President's Budget
 Exhibit R-1 FY 2011 Base and Overseas Contingency Operations (OCO) Request
 (Dollars in Thousands)

Appropriation: 0400D Research, Development, Test & Eval, DW

Date: February 2010

Line No	Program Element Number	Item	Act	FY 2009 (Base & OCO)	FY 2010 Base & OCO Enacted	FY 2010 Supplemental Request	FY 2010 Total	FY 2011 Base	FY 2011 OCO	FY 2011 Total Request	Section
6	0601384BP	Chemical and Biological Defense Program	01	60,355	78,764		78,764	49,508		49,508	U
	Basic Research			60,355	78,764		78,764	49,508		49,508	
14	0602384BP	Chemical and Biological Defense Program	02	231,331	224,830		224,830	169,287		169,287	U
	Applied Research			231,331	224,830		224,830	169,287		169,287	
34	0603384BP	Chemical and Biological Defense Program - Advanced Development	03	307,351	299,680		299,680	177,113		177,113	U
	Advanced Technology Development (ATD)			307,351	299,680		299,680	177,113		177,113	
78	0603884BP	Chemical and Biological Defense Program	04	69,793	209,275		209,275	277,062		277,062	U
	Advanced Component Development & Prototypes			69,793	209,275		209,275	277,062		277,062	
116	0604384BP	Chemical and Biological Defense Program	05	286,529	300,317		300,317	407,162		407,162	U
	System Development and Demonstration (SDD)			286,529	300,317		300,317	407,162		407,162	
148	0605384BP	Chemical and Biological Defense Program	06	100,470	106,033		106,033	120,995		120,995	U
149	0605502BP	Small Business Innovative Research - Chemical Biological Def	06	12,713							U
	RDT&E Management Support			113,183	106,033		106,033	120,995		120,995	
181	0607384BP	Chemical and Biological Defense (Operational Systems Development)	07	12,494	6,172		6,172	6,634		6,634	U
	Operational Systems Development			12,494	6,172		6,172	6,634		6,634	
Total Chemical and Biological Defense Program				1,081,036	1,225,071		1,225,071	1,207,761		1,207,761	

Exhibit R-1: FY 2011 President's Budget

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Exhibit R-2, RDT&E Budget Item Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 1: <i>Basic Research</i>	R-1 ITEM NOMENCLATURE PE 0601384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>
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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
Total Program Element	60.355	78.764	49.508	0.000	49.508	52.024	54.543	55.018	56.107	Continuing	Continuing
CB1: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	23.871	35.475	31.041	0.000	31.041	32.670	36.744	37.688	38.458	Continuing	Continuing
C11: <i>CONGRESSIONAL INTEREST ITEMS (BASIC RESEARCH)</i>	8.090	20.036	0.000	0.000	0.000	0.000	0.000	0.000	0.000	Continuing	Continuing
TB1: <i>MEDICAL BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	15.086	16.782	14.352	0.000	14.352	15.499	14.845	14.402	14.672	Continuing	Continuing
TC1: <i>MEDICAL CHEMICAL DEFENSE (BASIC RESEARCH)</i>	13.308	5.496	3.144	0.000	3.144	2.889	2.954	2.928	2.977	Continuing	Continuing
TR1: <i>MEDICAL RADIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	0.000	0.975	0.971	0.000	0.971	0.966	0.000	0.000	0.000	Continuing	Continuing

A. Mission Description and Budget Item Justification

This program element funds the Joint Service fundamental research program for (medical and physical sciences) Chemical, Biological, and Radiological (CBR) defense. The objective of the basic research program is to advance fundamental knowledge and understanding of the sciences with an emphasis in exploring new and innovative research for combating or countering chemical, biological and radiological weapons. Moreover, basic research supports a Joint Force concept of a lethal, integrated, supportable, highly mobile force with enhanced capability by the individual service member. Specifically, the program promotes theoretical and experimental research and studies in the chemical, biological, radiological, medical and related sciences. Research areas are aligned and prioritized to meet Joint Service needs as stated in mission area analyses, joint operational requirements and to take advantage of scientific opportunities. Basic research is executed by government laboratories, industry, and academia to include Historically Black Colleges and Universities and Minority Institutions (HBCU/MIs). Funds directed to these laboratories and research organizations capitalize on scientific talent, specialized facilities, and technological breakthroughs. The work in this program element is consistent with the Chemical Biological Defense Program Research, Development and Acquisition (RDA) Plan. Knowledge and technologies resulting from basic research efforts are expeditiously transitioned to the applied research (PE 0602384BP) and advanced technology development (PE 0603384BP) activities. This project also covers the conduct of basic research efforts in the areas of real-time sensing and immediate biological countermeasures. The projects in this PE are placed in BA1, because they are basic research efforts directed towards non-specific or non-unique military applications.

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Exhibit R-2, RDT&E Budget Item Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY	R-1 ITEM NOMENCLATURE
0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i>	PE 0601384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>
BA 1: <i>Basic Research</i>	

B. Program Change Summary (\$ in Millions)

	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011 Base</u>	<u>FY 2011 OCO</u>	<u>FY 2011 Total</u>
Previous President's Budget	61.194	58.974	0.000	0.000	0.000
Current President's Budget	60.355	78.764	49.508	0.000	49.508
Total Adjustments	-0.839	19.790	49.508	0.000	49.508
• Congressional General Reductions		-0.330			
• Congressional Directed Reductions		0.000			
• Congressional Rescissions	0.000	0.000			
• Congressional Adds		20.120			
• Congressional Directed Transfers		0.000			
• Reprogrammings	0.000	0.000			
• SBIR/STTR Transfer	-0.839	0.000			
• Other Adjustments	0.000	0.000	49.508	0.000	49.508

Congressional Add Details (\$ in Millions, and Includes General Reductions)

Project: CI1: *CONGRESSIONAL INTEREST ITEMS (BASIC RESEARCH)*

Congressional Add: *Garden State Cancer Center Vaccine Development Program -*

Congressional Add: *DNA Safeguard -*

Congressional Add: *In Vitro Models for Biodefense Vaccine -*

Congressional Add: *Superstructural Particle Evaluation and Characterization with Targeted Reaction Analysis (SPECTRA) -*

Congressional Add: *Defense Through Early Containment -*

Congressional Add: *Protection from Oxidative Stress -*

Congressional Add: *Research on a Molecular Approach to Hazardous Materials Decontamination -*

Congressional Add: *Synchotron Beamline and Experimental Station*

Congressional Add: *Advanced Development of Antiviral Prophylactics and Therapeutics*

Congressional Add: *Countermeasures to Chemical/Biological Control-Rapid Response*

Congressional Add: *MEMS Sensors for Real-Time Sensing of Weaponized Pathogens*

	<u>FY 2009</u>	<u>FY 2010</u>
	0.789	0.000
	1.184	0.000
	0.987	1.514
	1.184	0.000
	1.184	0.000
	1.579	0.000
	1.183	0.000
	0.000	3.187
	0.000	2.987
	0.000	2.788
	0.000	1.992

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R-1 Line Item #6

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Exhibit R-2, RDT&E Budget Item Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 1: <i>Basic Research</i>	R-1 ITEM NOMENCLATURE PE 0601384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>
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Congressional Add Details (\$ in Millions, and Includes General Reductions)

	FY 2009	FY 2010
Congressional Add: <i>Mismatch Repair Derived Antibody to Treat Staph Derived Bioweapon</i>	0.000	0.996
Congressional Add: <i>Portable Rapid Bacterial Warfare Detection</i>	0.000	3.983
Congressional Add: <i>Potent Human Monocolonal Antibodies Against BoNT, A, B and E Suited for Mass Production</i>	0.000	0.996
Congressional Add: <i>High Speed and High Volume Laboratory Network for Infectious Diseases</i>	0.000	1.593
Congressional Add Subtotals for Project: CI1	8.090	20.036
Congressional Add Totals for all Projects	8.090	20.036

Change Summary Explanation

Funding: N/A - Adjustments less than 10% of total program.

Schedule: N/A

Technical: N/A

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 1: <i>Basic Research</i>				R-1 ITEM NOMENCLATURE PE 0601384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>				PROJECT CB1: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>			
COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
CB1: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	23.871	35.475	31.041	0.000	31.041	32.670	36.744	37.688	38.458	Continuing	Continuing

A. Mission Description and Budget Item Justification

This project (CB1) supports basic research efforts in fundamental science phenomenology to include: life sciences; physical sciences; environmental sciences; mathematics; psychology and social sciences; and engineering. The objective of the Basic Research program is to successfully support the advancement of fundamental knowledge and understanding of the sciences with an emphasis on exploring new and innovative research for Chemical and Biological (CB) Defense. It includes new study areas, such as: nanoscale sciences; chemical, biological, and bio-inspired sciences; surface and signature sciences (with an emphasis on non-traditional agents (NTAs)); and information sciences. The aim is to promote innovative concepts and directions of research, which could lead to transformational capabilities to enhance the performance and ensure the safety of the Warfighter. Research in nanoscale sciences (nanoelectromechanical systems, molecular motors, and nanometer imaging) may bring about improvements in protection, decontamination and other core CB defense fields. Research in chemical, biological & bio-inspired sciences includes research in concepts such as synthetic biology, biomimetics, and other emerging areas of science to build a foundation for developing novel smart materials. This will combine multiple functionalities into a common autonomous unit or network. Surface and signature sciences focuses on the study of physical and chemical properties, especially with regard to NTAs, that seek to improve physical capabilities such as detection and decontamination. Informational Sciences includes research in understanding cognitive and physiological effects on human decision-making, behavior and performance, and modeling and simulation of CB threats. Breakthroughs and advances in functional capabilities gained from these scientific disciplines could impact the entire chemical and biological defense science and technology program. Basic research activities described in this budget justification leverage existing research programs and activities within the DoD and other government agencies to accelerate transformational breakthroughs, which may be transitioned to applied research or advanced development initiatives. Due to the exploratory, academic, and theoretical nature of basic research efforts, projects described in this justification typically have a duration period, from conception to completion, of three to five years. Promising basic research efforts will be further exploited for their application to chemical and biological defense in Budget Activity 2 (Applied Science). The basic research program promotes cross-pollination between government and academia, as well as sponsors world class scientists while promoting the development of young researchers.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) SBIR	0.000	0.584	0.000	0.000	0.000

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 1: <i>Basic Research</i>	R-1 ITEM NOMENCLATURE PE 0601384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	PROJECT CB1: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>				
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2010 Plans:</i> Small Business Innovative Research.						
2) Basic Research Core Nano-Scale Sciences: Improve understanding of nano-scale materials (scale of 1-100 nanometers in length) for use in chemical and biological defense. <i>FY 2009 Accomplishments:</i> Completed efforts investigating new types of materials for potential use in decontamination and protection, and shared information on new techniques for detection of chemical agents through novel applications of physics and chemistry. Continued study of compounds which mimic biological organisms and nano-scale sensing technologies for identification of agents. Continued studies of new materials being developed through nanotechnology for protective equipment, while initiating new efforts into new textiles with a higher resistance to oily substances or with adjustable porosity. Other new efforts studied interfaces between nano-materials and living cells, and systems found in nature for creative solutions for future protection concepts. <i>FY 2010 Plans:</i> Complete study of some compounds which mimic biological organisms and nano-scale sensing technologies for identification of agents. Continue efforts into new textiles with a higher resistance to oily substances or with adjustable porosity, as well, as efforts studying interfaces between nano-materials and living cells. Continue the study of systems found in nature for creative solutions for future protection concepts. Continue to identify new topics for investments in basic research to support the fundamental scientific phenomena in nano-scale science technology. Investigate new concepts in nano-scale chemical and biological sensing/detection. Initiate new studies to develop nano-scaled porous materials. Identify/leverage state-of-the-art breakthroughs to fill capability gaps.		5.572	9.009	8.700	0.000	8.700

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 1: <i>Basic Research</i>	R-1 ITEM NOMENCLATURE PE 0601384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	PROJECT CB1: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>				
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p>Begin developing novel tools to investigate cells and cell mechanisms. Characterize NTA toxicokinetic properties and mechanisms of toxicity for NTAs. Assess effectiveness of developmental general purpose decontaminants, as well as, explore new formulations. Maintain visibility of relevant research which could be leveraged for the benefit of chemical and biological defense.</p> <p><i>FY 2011 Base Plans:</i> Continue developing novel tools to investigate cells and cell mechanisms. Continue to investigate and leverage developments in bioscience, bio-inspired science, and chemical sciences to support and improve fundamental scientific understanding. Leverage and merge developments with other basic research areas such as information sciences and surface and signature sciences. Initiate efforts in response to identified science gaps.</p>						
<p>4) Basic Research Core</p> <p>Information Science: Leverages new developments in information and computation to impact modeling and other chemical and biological defense efforts.</p> <p><i>FY 2009 Accomplishments:</i> Continued research on projects initiated in FY08. Initiated efforts to investigate genetic algorithms to identify optimal material arrangements, quantification and reduction of uncertainty for dispersion models via meteorological predictions through computer experimentation, calculations of the complete electromagnetic response of large macromolecules, and new molecular recognition signatures in the electromagnetic spectrum.</p> <p><i>FY 2010 Plans:</i> Continue FY08/FY09 projects. Initiate efforts to support and investigate genetic algorithms. Seek to understand cognitive effects of heightened sensory input. Research conducted will draw from many disciplines, including: cognitive psychology; neuroscience; linguistics; medical sciences; and</p>		5.925	5.876	6.000	0.000	6.000

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 1: <i>Basic Research</i>	R-1 ITEM NOMENCLATURE PE 0601384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	PROJECT CB1: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>				
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
will leverage advances in physics, mathematics, biology, and other relevant sciences to improve informational and decision making tools. <i>FY 2011 Base Plans:</i> Continue investigating genetic algorithms and studying effects of heightened sensory input during chemical biological warfare events. Utilize efforts in information sciences to inform other areas of core chemical and biological defense programs, such as modeling and computational efforts.						
5) Basic Research Core Cognitive Science: Focuses on thinking and decision making to impact support tools for CB defense. <i>FY 2009 Accomplishments:</i> Continued research on projects initiated in FY08. Initiated efforts to investigate the presentation of risk and uncertainty for chemical and biological defense decision making. <i>FY 2010 Plans:</i> All Cognitive Science efforts are re-aligned to Information Science.		3.463	0.000	0.000	0.000	0.000
6) Basic Research Core Integration of Basic Research Science: Focuses on basic research for chemical and biological defense and reaches out to a varied performer base for the best innovations and programs. <i>FY 2009 Accomplishments:</i> Completed research on projects initiated in FY08, and transitioned relevant information to various physical applied research projects located in Budget Activity 2.		4.111	0.000	0.000	0.000	0.000
7) Basic Research Core		0.000	8.488	8.000	0.000	8.000

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 1: <i>Basic Research</i>	R-1 ITEM NOMENCLATURE PE 0601384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	PROJECT CB1: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>
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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p>Surface and Signature Sciences: A new study area established to focus on the study of physical and chemical properties that seeks to improve physical capabilities, such as, detection and decontamination.</p> <p><i>FY 2010 Plans:</i> Identify and exploit novel tools to investigate surface and signature sciences to inform capability gaps in fields such as detection and decontamination. Initiate and combine the efforts that improve the phenomenology needed to protect, detect, decontaminate, or otherwise counter chemical (to include NTAs) and biological threats. Study interactions of chemical and biological agents with biological and environmental matrices.</p> <p><i>FY 2011 Base Plans:</i> Continue studying interactions of chemical and biological agents with biological and environmental matrices, and develop novel tools to investigate surface and signature sciences to address capability gaps. Study signature sciences and surface interactions.</p>					
Accomplishments/Planned Programs Subtotals	23.871	35.475	31.041	0.000	31.041

C. Other Program Funding Summary (\$ in Millions)

Line Item	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	FY 2012	FY 2013	FY 2014	FY 2015	Cost To Complete	Total Cost
• CB2: <i>CHEMICAL BIOLOGICAL DEFENSE (APPLIED RESEARCH)</i>	102.599	110.955	88.897		88.897	100.243	97.979	90.686	91.554	Continuing	Continuing
• CB3: <i>CHEMICAL BIOLOGICAL DEFENSE (ATD)</i>	19.567	25.297	15.410		15.410	21.450	26.120	36.775	37.148	Continuing	Continuing

D. Acquisition Strategy

N/A

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY	R-1 ITEM NOMENCLATURE	PROJECT
0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 1: <i>Basic Research</i>	PE 0601384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	CB1: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>

E. Performance Metrics

N/A

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 1: <i>Basic Research</i>	R-1 ITEM NOMENCLATURE PE 0601384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	PROJECT C11: <i>CONGRESSIONAL INTEREST ITEMS (BASIC RESEARCH)</i>
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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
C11: <i>CONGRESSIONAL INTEREST ITEMS (BASIC RESEARCH)</i>	8.090	20.036	0.000	0.000	0.000	0.000	0.000	0.000	0.000	Continuing	Continuing

A. Mission Description and Budget Item Justification

The efforts listed in this project include congressional interest programs for FY09 and FY10.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010
Congressional Add: Garden State Cancer Center Vaccine Development Program - <i>FY 2009 Accomplishments:</i> Continued the development of a safe vaccine against smallpox that does not require whole or live virus, thereby eliminating the danger of vaccine-associated side effects and transmission for viral infections to immunocompromised individuals.	0.789	0.000
Congressional Add: DNA Safeguard - <i>FY 2009 Accomplishments:</i> Continued development of a stable, DNA-based chemical marker (DNA Barcode) capable of encoding information that can be added to any DNA sample in order to label the sample and guarantee its integrity.	1.184	0.000
Congressional Add: In Vitro Models for Biodefense Vaccine -	0.987	1.514

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 1: <i>Basic Research</i>	R-1 ITEM NOMENCLATURE PE 0601384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	PROJECT C11: <i>CONGRESSIONAL INTEREST ITEMS (BASIC RESEARCH)</i>
B. Accomplishments/Planned Program (\$ in Millions)		
	FY 2009	FY 2010
<i>FY 2009 Accomplishments:</i> Conducted basic research for the use of In Vitro models in vaccine development.		
<i>FY 2010 Plans:</i> Continuation of FY09 research		
Congressional Add: Superstructural Particle Evaluation and Characterization with Targeted Reaction Analysis (SPECTRA) - <i>FY 2009 Accomplishments:</i> Continued basic research on superstructural particle evaluation and characterization with targeted reaction analysis begun in FY06.	1.184	0.000
Congressional Add: Defense Through Early Containment - <i>FY 2009 Accomplishments:</i> Conducted basic research focused on containment of agents following an incident.	1.184	0.000
Congressional Add: Protection from Oxidative Stress - <i>FY 2009 Accomplishments:</i> Conducted basic research focused on protection technologies.	1.579	0.000
Congressional Add: Research on a Molecular Approach to Hazardous Materials Decontamination - <i>FY 2009 Accomplishments:</i> Continued research on molecular approach to decontamination in collaboration with the Naval Surface Warfare Center (NSWC) begun in FY06.	1.183	0.000

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program		DATE: February 2010	
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 1: <i>Basic Research</i>	R-1 ITEM NOMENCLATURE PE 0601384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	PROJECT C11: <i>CONGRESSIONAL INTEREST ITEMS (BASIC RESEARCH)</i>	
B. Accomplishments/Planned Program (\$ in Millions)			
		FY 2009	FY 2010
Congressional Add: Synchrotron Beamline and Experimental Station <i>FY 2010 Plans:</i> Build an experimental end-station at National Synchrotron Light Source-II for the purpose of conducting basic research on the structure and processes of pathogens, toxins and their antidotes.		0.000	3.187
Congressional Add: Advanced Development of Antiviral Prophylactics and Therapeutics <i>FY 2010 Plans:</i> Apply knowledge of the discovery of a new class of antiviral drugs to develop medical countermeasures against viral biowarfare agents and for therapies against other viral diseases such as Hepatitis C, HIV and Influenza. Explore chemical modifications of previously identified broadly active compounds to target proteins involved in additional viruses of interest. Demonstrate efficacy of active compounds via in vitro and animal trials.		0.000	2.987
Congressional Add: Countermeasures to Chemical/Biological Control-Rapid Response <i>FY 2010 Plans:</i> Research Support of Biodefense and emerging infectious disease.		0.000	2.788
Congressional Add: MEMS Sensors for Real-Time Sensing of Weaponized Pathogens <i>FY 2010 Plans:</i> Develop a wearable sensor to detect weaponized pathogens utilizing the unique properties of diamond and enable a new class of compact, wearable chemical and biological point sensors, with unprecedented sensitivity, stability, and reproducibility.		0.000	1.992
		0.000	0.996

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 1: <i>Basic Research</i>	R-1 ITEM NOMENCLATURE PE 0601384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	PROJECT C11: <i>CONGRESSIONAL INTEREST ITEMS (BASIC RESEARCH)</i>
B. Accomplishments/Planned Program (\$ in Millions)		
	FY 2009	FY 2010
Congressional Add: Mismatch Repair Derived Antibody to Treat Staph Derived Bioweapon <i>FY 2010 Plans:</i> Develop fully human anti-Staphylococcus enterotoxin B (SEB) monoclonal antibodies (mAbs) that can neutralize >1000 times the human LD50 of the toxin.		
Congressional Add: Portable Rapid Bacterial Warfare Detection <i>FY 2010 Plans:</i> Develop a field deployable system based on InfraRed spectroscopy.	0.000	3.983
Congressional Add: Potent Human Monocolonal Antibodies Against BoNT, A, B and E Suited for Mass Production <i>FY 2010 Plans:</i> Develop Potent Human Monocolonal Antibodies Against BoNT, A, B and E Suited for Mass Production	0.000	0.996
Congressional Add: High Speed and High Volume Laboratory Network for Infectious Diseases <i>FY 2010 Plans:</i> Develop an expanded capability to include other biothreat agents, including bacterial and/or viruses (dual-use).	0.000	1.593
Congressional Adds Subtotals	8.090	20.036

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 1: <i>Basic Research</i>	R-1 ITEM NOMENCLATURE PE 0601384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	PROJECT C11: <i>CONGRESSIONAL INTEREST ITEMS (BASIC RESEARCH)</i>
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C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011</u> <u>Base</u>	<u>FY 2011</u> <u>OCO</u>	<u>FY 2011</u> <u>Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To</u> <u>Complete</u>	<u>Total Cost</u>
• C12: <i>CONGRESSIONAL INTEREST ITEMS (APPLIED RESEARCH)</i>	42.714	16.630	0.000		0.000	0.000	0.000	0.000	0.000	Continuing	Continuing
• C13: <i>CONGRESSIONAL INTEREST ITEMS (ATD)</i>	46.971	18.622	0.000		0.000	0.000	0.000	0.000	0.000	Continuing	Continuing

D. Acquisition Strategy

N/A

E. Performance Metrics

N/A

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program								DATE: February 2010			
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 1: <i>Basic Research</i>				R-1 ITEM NOMENCLATURE PE 0601384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>				PROJECT TB1: <i>MEDICAL BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>			
COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
TB1: <i>MEDICAL BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	15.086	16.782	14.352	0.000	14.352	15.499	14.845	14.402	14.672	Continuing	Continuing

A. Mission Description and Budget Item Justification

This project (TB1) funds basic research of vaccines, diagnostic tools, and therapeutic drugs to provide effective medical defense against validated biological threat agents including bacteria, toxins, and viruses. Advance innovative biotechnology approaches with the potential to rapidly identify, diagnose, prevent, and treat disease due to exposure to biological threat agents. Project categories include core science efforts and technology programs areas in biological defense capability areas, such as Pretreatments, Diagnostics, and Therapeutics. Starting in FY10, all efforts will be combined into a single capability area called Biological Based Basic Research.

This project also includes efforts such as the Transformational Medical Technologies Initiative (TMTI). The TMTI was launched to respond to the threat of emerging or intentionally bioengineered biological threats. TMTI's mission is to protect the Warfighter from genetically engineered biological threats by providing a rapid response capability from identification of pathogens to the delivery of medical countermeasures. This mission is accomplished through two main efforts: 1) developing broad spectrum (multi-agent) therapeutics against biological warfare (BW) agents (e.g, one drug that treats multiple agents); and 2) developing platform technologies to assist in the rapid development of medical countermeasures (MCMs) in response to BW agents (e.g, developing new and innovative ways to mass produce drugs in the event of a biological incident).

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.	0.000	0.270	0.000	0.000	0.000
2) Diagnostics Pursue technologies that enable medical elements to determine exposure or infection of forces by a biological warfare agent and assist in appropriate lifesaving treatment.	3.026	0.000	0.000	0.000	0.000

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 1: <i>Basic Research</i>	R-1 ITEM NOMENCLATURE PE 0601384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	PROJECT TB1: <i>MEDICAL BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>				
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2009 Accomplishments:</i> Continued to seek novel avenues for assay design and application. Investigated cutting edge technologies as new genomic techniques become available. Accelerated identification of novel biomarkers of biological warfare agent (BWA) infection and applied to assay development.</p> <p><i>FY 2010 Plans:</i> Efforts re-aligned to Biological Based Basic Research.</p>						
<p>3) Pretreatments</p> <p>Multiagent Vaccines: Research vaccines that protect against multiple agents.</p> <p><i>FY 2009 Accomplishments:</i> Utilized novel technologies to define target antigens for different bio-threat pathogens. Explored DNA-based vaccine formulations against multiple agents. Incorporated novel adjuvants and/or delivery systems in the design of a multi-agent vaccine.</p> <p><i>FY 2010 Plans:</i> Efforts re-aligned to Biological Based Basic Research.</p>		0.315	0.000	0.000	0.000	0.000
<p>4) Pretreatments</p> <p>Vaccine Research Support: Research human immune response and pathogenicity of biological agents.</p> <p><i>FY 2009 Accomplishments:</i> Conducted basic pathogenicity studies of selected biothreat agents. Developed and refined in vitro correlates of immunity for new antigen in relation to vaccines under development. Pursued the identification and evaluation of novel target antigens for intracellular pathogens by studying the innate and adaptive immune responses to pathogens. Optimized epitope mapping of lead antigen candidates.</p>		2.937	0.000	0.000	0.000	0.000

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 1: <i>Basic Research</i>	R-1 ITEM NOMENCLATURE PE 0601384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	PROJECT TB1: <i>MEDICAL BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>				
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2010 Plans:</i> Efforts re-aligned to Biological Based Basic Research.						
5) Therapeutics Viral Therapeutics: Research understanding of viral infection. <i>FY 2009 Accomplishments:</i> Delineated the mechanisms of pathogenesis of conventional threats to support the progression of therapeutics to advanced development. Compared the host response of well characterized threats with that of poorly characterized category A and B threats to identify new therapeutic targets. <i>FY 2010 Plans:</i> Effort re-aligned to Biological Based Basic Research.		0.435	0.000	0.000	0.000	0.000
6) Therapeutics Toxin Therapeutics: Research efforts to enhance understanding of toxins and their effects on the host. <i>FY 2009 Accomplishments:</i> Improved in silico, in vitro, and in vivo modeling systems that will assist in defining responses to threat agent toxins. Completed development of a mouse model for inhalational exposure to staphylococcal enterotoxin B (SEB) using microinstillation technology. Characterized the process of intracellular targeting of BoNT, and initiated intracellular assay model development. Defined the cellular factors responsible for Botulinum Neurotoxin (BoNT) translocation inside cells. Determined the structural requirements of potential restorative therapeutics for neuromuscular paralysis following BoNT intoxication. <i>FY 2010 Plans:</i> Efforts re-aligned to Biological Based Basic Research.		2.606	0.000	0.000	0.000	0.000

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 1: <i>Basic Research</i>	R-1 ITEM NOMENCLATURE PE 0601384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	PROJECT TB1: <i>MEDICAL BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>				
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
7) Therapeutics Bacterial Therapeutics: Research efforts to enhance understanding of bacterial pathogens. <i>FY 2009 Accomplishments:</i> Characterized new potential targets for therapeutic countermeasures, focusing on those identified for poorly characterized threats. <i>FY 2010 Plans:</i> Efforts re-aligned to Biological Based Basic Research.		0.653	0.000	0.000	0.000	0.000
8) Biological Based Basic Research Research to understand biological agents of interest, their pathways, virulence, immunization factors and identification. <i>FY 2010 Plans:</i> Determine mechanisms of pathogenesis for viral and bacterial biothreat agents and toxins. Define immune responses and mechanisms that confer protection against biothreat agents. Identify novel and/or shared antigens from viral and bacterial threat agents to be used in the design of future vaccine formulations. Determine the contribution of post-translational modification of Botulinum Neurotoxin (BoNT) to the intracellular biology of the toxin. Determine advanced pharmacokinetic models of BoNT intoxication to define the therapeutic window of opportunity. <i>FY 2011 Base Plans:</i> Conduct studies of pathogenic mechanisms for viral and bacterial biothreat agents and toxins. Clarify mechanisms of host-pathogen interaction to identify mechanisms of pathogenesis and/or correlates of protective immunity against biothreat agents. Define novel and/or shared antigens from viral and bacterial threat agents to be used in the design of future treatment options. Define the contribution		0.000	9.160	8.899	0.000	8.899

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010	
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 1: <i>Basic Research</i>		R-1 ITEM NOMENCLATURE PE 0601384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>		PROJECT TB1: <i>MEDICAL BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	
B. Accomplishments/Planned Program (\$ in Millions)					
of post-translational modification to the structure and biology of BoNT. Research novel constructs for affinity reagents for the identification of biological warfare agents and biomarkers.					
9) Transformational Medical Technologies Initiative					
<p>Multiagent (Broad Spectrum) Medical Countermeasures: Basic research efforts focused on the early drug discovery phase of drug development. Active monitoring of scientific literature to generate hypotheses for research. Review scientific findings and assess a foundation for characterizing new therapeutics. Identify and develop brand new compounds that could lead to successful therapeutic candidates. Scientific studies to generate research ideas, hypotheses, and experimental designs for addressing the development of therapeutics against Biological Warfare (BW) agents. Use computer simulation or other virtual platforms to test hypotheses. Begin research, data collection, and analysis to test hypothesis. Explore alternative concepts, identify and evaluate critical technologies and components, and begin characterization of candidates. Demonstrate preliminary efficacy.</p> <p><i>FY 2009 Accomplishments:</i> Continued drug discovery research for broad-spectrum countermeasures with new candidates. Continued basic research to identify new candidates for molecular targets for broad-spectrum countermeasures. Continued to evaluate new thrust areas in genomics, proteomics, bioinformatics, and other relevant systems biology research. Focused efforts on promising intervention points for broad-spectrum therapeutic approaches based on results from drug design collaborations. Developed computer models and other methodologies to support rational drug design by determining the three-dimensional structure of important molecules based on the genetic sequences of organisms. Continued to study changes in host response to infection. Initiated study of biomarkers for intracellular bacterial (ICB) and hemorrhagic fever virus (HFV) agents.</p> <p><i>FY 2010 Plans:</i> Initiate support for the discovery of conserved host and pathogen directed targets for the development of broad spectrum drugs against BW agents. Validate computer models and other methodologies for</p>					
	5.114	5.471	0.000	0.000	0.000

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 1: <i>Basic Research</i>	R-1 ITEM NOMENCLATURE PE 0601384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	PROJECT TB1: <i>MEDICAL BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>
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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
technologies are appropriate for each aspect of the countermeasure development. Continue to support discovery of conserved host and pathogen directed targets for the development of broad spectrum drugs against BW agents. Continue to develop leading edge technologies to assist in pathogen characterization, target identification, countermeasure discovery and countermeasure evaluation.					
Accomplishments/Planned Programs Subtotals	15.086	16.782	14.352	0.000	14.352

C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011 Base</u>	<u>FY 2011 OCO</u>	<u>FY 2011 Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To Complete</u>	<u>Total Cost</u>
• TB2: <i>MEDICAL BIOLOGICAL DEFENSE (APPLIED RESEARCH)</i>	50.485	53.930	43.858		43.858	50.866	51.077	51.051	51.959	Continuing	Continuing
• TB3: <i>MEDICAL BIOLOGICAL DEFENSE (ATD)</i>	180.425	203.723	115.233		115.233	125.666	109.737	115.049	117.289	Continuing	Continuing

D. Acquisition Strategy

N/A

E. Performance Metrics

N/A

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY			R-1 ITEM NOMENCLATURE				PROJECT				
0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 1: <i>Basic Research</i>			PE 0601384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>				TC1: <i>MEDICAL CHEMICAL DEFENSE (BASIC RESEARCH)</i>				
COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
TC1: <i>MEDICAL CHEMICAL DEFENSE (BASIC RESEARCH)</i>	13.308	5.496	3.144	0.000	3.144	2.889	2.954	2.928	2.977	Continuing	Continuing

A. Mission Description and Budget Item Justification

This project (TC1) emphasizes the understanding of the basic action mechanisms of nerve, blister, blood, and respiratory agents within the body. Basic studies are performed to delineate biological mechanisms for identified and emerging chemical threats to generate required information for initial design and synthesis of chemical medical countermeasures. In addition, these studies are further designed to maintain and extend a science base. Starting in FY10, all efforts will be combined into a new capability area termed Chemical Based Basic Research.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.	0.000	0.093	0.000	0.000	0.000
2) Therapeutics Respiratory and Systemic: Research efforts that define pathways of injury and therapeutic targets against chemical agent exposure through inhalation. <i>FY 2009 Accomplishments:</i> Expanded efforts to elucidate common injury pathways due to multiple agents and routes of exposure, to maximize application to the development of broad-based therapeutics. Established definitive correlation between simulants and live agent effects at the molecular level. <i>FY 2010 Plans:</i> Efforts re-aligned to Chemical Based Basic Research.	5.133	0.000	0.000	0.000	0.000

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 1: <i>Basic Research</i>	R-1 ITEM NOMENCLATURE PE 0601384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	PROJECT TC1: <i>MEDICAL CHEMICAL DEFENSE (BASIC RESEARCH)</i>				
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
3) Therapeutics Cutaneous and Ocular: Research efforts that define pathways of injury and therapeutic targets for chemical agent exposure through skin and eye exposure. <i>FY 2009 Accomplishments:</i> Extrapolated the results of genotoxicity studies to the development of cancerous conditions using the appropriate in vivo models. Investigated the effects of solvent vehicles on percutaneous transmission to normalize past, present, and future research endeavors. Investigated new tissue engineering technologies to reduce reliance on grafts. <i>FY 2010 Plans:</i> Efforts re-aligned to Chemical Based Basic Research.		2.541	0.000	0.000	0.000	0.000
4) Therapeutics Neurologic: Research efforts that aim to improve understanding of nerve agents. <i>FY 2009 Accomplishments:</i> Researched mechanisms of action of nerve agents and therapeutic interventions using whole animal models, with a focus on data required to support FDA submissions. Initiated research into the development of nerve agent therapeutic alternatives with reduced impact on visual performance. <i>FY 2010 Plans:</i> Efforts re-aligned to Chemical Based Basic Research.		1.703	0.000	0.000	0.000	0.000
5) Therapeutics Medical Toxicology: Research Non Traditional Agents (NTAs) and other agents to improve understanding of NTA exposure.		3.931	0.000	0.000	0.000	0.000

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 1: <i>Basic Research</i>	R-1 ITEM NOMENCLATURE PE 0601384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	PROJECT TC1: <i>MEDICAL CHEMICAL DEFENSE (BASIC RESEARCH)</i>				
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2009 Accomplishments:</i> Demonstrated the biological equivalency of NTA toxicity mechanisms across relevant species.</p> <p><i>FY 2010 Plans:</i> Efforts re-aligned to Chemical Based Basic Research.</p>						
<p>6) Chemical Based Basic Research (CBBR)</p> <p>Research focuses on understanding chemical agents, their mechanism of action, toxicity, cellular injury, and identification.</p> <p><i>FY 2010 Plans:</i> Investigate new tissue engineering technologies to reduce reliance on skin grafts. Assess the results of genotoxicity studies. Research mechanisms of action of nerve agents and therapeutic interventions using whole animal models, with a focus on data required to support FDA submissions. Initiate research into the development for novel nerve agent therapeutics with reduced impact on visual performance. Initiate development of new animal models to characterize in vivo effects of NTAs. Demonstrate the biological equivalency of Non-Traditional Agent (NTA) toxicity mechanisms across relevant species.</p> <p><i>FY 2011 Base Plans:</i> Research pathways of molecular mechanisms of injury associated with chemical warfare agents. Conduct mechanistic studies using appropriate in vitro models to identify the biochemical cascade of effects following chemical agent exposure. Based on these studies, generate basic information for initial design and synthesis of medical countermeasures, located in Budget Activity 2, Project TC2.</p>		0.000	5.403	3.144	0.000	3.144
Accomplishments/Planned Programs Subtotals		13.308	5.496	3.144	0.000	3.144

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 1: <i>Basic Research</i>	R-1 ITEM NOMENCLATURE PE 0601384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	PROJECT TC1: <i>MEDICAL CHEMICAL DEFENSE (BASIC RESEARCH)</i>

C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011</u> <u>Base</u>	<u>FY 2011</u> <u>OCO</u>	<u>FY 2011</u> <u>Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To</u> <u>Complete</u>	<u>Total Cost</u>
• TC2: <i>MEDICAL CHEMICAL DEFENSE (APPLIED RESEARCH)</i>	35.008	40.418	33.648		33.648	36.327	36.500	37.475	38.150	Continuing	Continuing
• TC3: <i>MEDICAL CHEMICAL DEFENSE (ATD)</i>	21.641	28.971	29.134		29.134	30.401	30.546	31.356	31.877	Continuing	Continuing

D. Acquisition Strategy

N/A

E. Performance Metrics

N/A

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program								DATE: February 2010			
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 1: <i>Basic Research</i>				R-1 ITEM NOMENCLATURE PE 0601384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>				PROJECT TR1: <i>MEDICAL RADIOLOGICAL DEFENSE (BASIC RESEARCH)</i>			
COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
TR1: <i>MEDICAL RADIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	0.000	0.975	0.971	0.000	0.971	0.966	0.000	0.000	0.000	Continuing	Continuing

A. Mission Description and Budget Item Justification

This project (TR1) emphasizes the research and study of medical countermeasures to protect the Warfighter against radiation exposure. Specifically, this project identifies the basic action mechanisms of Acute Radiation Syndrome (ARS) and Delayed Effects of Acute Radiation Exposure (DEARE), as well as, develops possible radioprotectants (Pretreatments), post-irradiation exposure treatments (Therapeutics), and the ability to identify exposure to radiation (Diagnostics). These Basic Research efforts advance promising technology with the potential to rapidly identify, diagnose, prevent, and mitigate ARS and/or DEARE in the event of a radiological incident.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) Medical Radiological Defense Research focuses on understanding mechanisms of injury from radiation exposure. <i>FY 2010 Plans:</i> Initiate efforts to identify mechanisms of injury from acute radiation exposure and delayed health effects following radiation exposure. Explore novel assays to diagnose radiation injury, through studies of cellular science, metabolism, and bioregulators. <i>FY 2011 Base Plans:</i> Continue projects begun in FY10 to understand cellular and molecular responses to ionizing radiation and identify biomarkers of radiation exposure.	0.000	0.959	0.971	0.000	0.971
2) SBIR	0.000	0.016	0.000	0.000	0.000

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 1: <i>Basic Research</i>	R-1 ITEM NOMENCLATURE PE 0601384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	PROJECT TR1: <i>MEDICAL RADIOLOGICAL DEFENSE (BASIC RESEARCH)</i>
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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2010 Plans:</i> Small Business Innovative Research.					
Accomplishments/Planned Programs Subtotals	0.000	0.975	0.971	0.000	0.971

C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011 Base</u>	<u>FY 2011 OCO</u>	<u>FY 2011 Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To Complete</u>	<u>Total Cost</u>
• TR2: <i>MEDICAL RADIOLOGICAL DEFENSE (APPLIED RESEARCH)</i>	0.525	2.897	2.884		2.884	1.904	2.855	1.913	1.903	Continuing	Continuing
• TR3: <i>MEDICAL RADIOLOGICAL DEFENSE (ATD)</i>	4.859	2.403	0.957		0.957	0.966	1.922	2.901	2.927	Continuing	Continuing

D. Acquisition Strategy

N/A

E. Performance Metrics

N/A

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Exhibit R-2, RDT&E Budget Item Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 2: <i>Applied Research</i>	R-1 ITEM NOMENCLATURE PE 0602384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (APPLIED RESEARCH)</i>
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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
Total Program Element	231.331	224.830	169.287	0.000	169.287	189.340	188.411	181.125	183.566	Continuing	Continuing
CB2: <i>CHEMICAL BIOLOGICAL DEFENSE (APPLIED RESEARCH)</i>	102.599	110.955	88.897	0.000	88.897	100.243	97.979	90.686	91.554	Continuing	Continuing
CI2: <i>CONGRESSIONAL INTEREST ITEMS (APPLIED RESEARCH)</i>	42.714	16.630	0.000	0.000	0.000	0.000	0.000	0.000	0.000	Continuing	Continuing
TB2: <i>MEDICAL BIOLOGICAL DEFENSE (APPLIED RESEARCH)</i>	50.485	53.930	43.858	0.000	43.858	50.866	51.077	51.051	51.959	Continuing	Continuing
TC2: <i>MEDICAL CHEMICAL DEFENSE (APPLIED RESEARCH)</i>	35.008	40.418	33.648	0.000	33.648	36.327	36.500	37.475	38.150	Continuing	Continuing
TR2: <i>MEDICAL RADIOLOGICAL DEFENSE (APPLIED RESEARCH)</i>	0.525	2.897	2.884	0.000	2.884	1.904	2.855	1.913	1.903	Continuing	Continuing

A. Mission Description and Budget Item Justification

Funding under this program element (PE) sustains a robust defense program, which both reduces the danger of a chemical, biological, or radiological (CBR) attack and enables U.S. forces to survive, and continue operations in a CBR environment. The medical program focuses on development of antidotes, drug treatments, casualty diagnosis, patient decontamination and medical technologies management. In the physical sciences area, the emphasis is on continuing improvements in CB defense materiel, including contamination avoidance, decontamination, and protection technologies. Research efforts are planned to be initiated for CB defense technologies that will result from a strategic approach of converging nanotechnology, biotechnology, information technology and cognitive science. This PE also provides for applied research in the areas of real-time sensing and immediate biological countermeasures. The work in this PE is consistent with the Chemical Biological Defense Program Research Development and Acquisition (RDA) Plan. Efforts under this PE transition to or provide risk reduction for Advanced Technology Development (PE: 0603384BP), Advanced Component Development and Prototypes (PE: 0603884BP) and System Development and Demonstration (PE: 0604384BP). This project is placed in BA2, because it includes non-system specific development, directed toward military needs.

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Exhibit R-2, RDT&E Budget Item Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY	R-1 ITEM NOMENCLATURE
0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i>	PE 0602384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (APPLIED RESEARCH)</i>
BA 2: <i>Applied Research</i>	

B. Program Change Summary (\$ in Millions)

	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011 Base</u>	<u>FY 2011 OCO</u>	<u>FY 2011 Total</u>
Previous President's Budget	239.297	209.072	0.000	0.000	0.000
Current President's Budget	231.331	224.830	169.287	0.000	169.287
Total Adjustments	-7.966	15.758	169.287	0.000	169.287
• Congressional General Reductions		-0.942			
• Congressional Directed Reductions		0.000			
• Congressional Rescissions	0.000	0.000			
• Congressional Adds		16.700			
• Congressional Directed Transfers		0.000			
• Reprogrammings	4.731	0.000			
• SBIR/STTR Transfer	-2.697	0.000			
• Other Adjustments	-10.000	0.000	169.287	0.000	169.287

Congressional Add Details (\$ in Millions, and Includes General Reductions)

Project: CI2: *CONGRESSIONAL INTEREST ITEMS (APPLIED RESEARCH)*

Congressional Add: *Rapid Forensic Evaluation of Microbes in Biodefense*

Congressional Add: *Chem/Bio IR Detection System*

Congressional Add: *Zumwalt National Program for Countermeasures to Bio Chem Threats*

Congressional Add: *HyperAcute Vaccine Development*

Congressional Add: *Antibody-based Therapeutic against Smallpox*

Congressional Add: *Novel Viral Biowarfare Agent Identification and Treatment (NOVBAIT)*

Congressional Add: *Mixed Oxidants for Chemical and Biological Decontamination*

Congressional Add: *Bio Surety Development and Management Program -*

Congressional Add: *Countermeasures to Chemical/Biological Control-Rapid Response -*

Congressional Add: *Multiple Applications for Light Activated, Reactive Materials for Protection of Warfighter, First Responder, and Public Health -*

	<u>FY 2009</u>	<u>FY 2010</u>
	0.989	0.000
	1.186	1.892
	1.187	0.000
	2.373	3.585
	0.791	0.000
	3.955	0.000
	2.769	0.000
	1.186	0.000
	2.372	0.000
	1.582	0.000

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Exhibit R-2, RDT&E Budget Item Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 2: <i>Applied Research</i>	R-1 ITEM NOMENCLATURE PE 0602384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (APPLIED RESEARCH)</i>
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Congressional Add Details (\$ in Millions, and Includes General Reductions)	FY 2009	FY 2010
Congressional Add: <i>Chemical Biological Preparedness Center for Advanced Development of Mobile Rapid Response Prototype -</i>	3.955	0.000
Congressional Add: <i>Novel System for Developing Therapeutics against Botulism -</i>	3.955	0.000
Congressional Add: <i>Ultra-Rapid Next Generation Pathogen Identification -</i>	1.978	0.000
Congressional Add: <i>Preventing Long-Term Brain and Lung Damage Caused by Battlefield Trauma Project -</i>	2.868	0.000
Congressional Add: <i>Chemical Agent Fate Appropriate Response Tool -</i>	1.582	1.593
Congressional Add: <i>Multivalent Marburg/Ebola Vaccine -</i>	3.461	0.000
Congressional Add: <i>Botulinum Neurotoxin Research -</i>	1.582	1.992
Congressional Add: <i>Miniaturized Chemical Detector for Chemical Warfare Protection (ChemPen) -</i>	1.581	1.593
Congressional Add: <i>Continued Expansion of Prototypes for Destruction of Airborne Pathogen -</i>	0.791	0.000
Congressional Add: <i>Mismatch Repair Derived Antibody to Treat Staph Derived Bioweapon -</i>	1.582	0.000
Congressional Add: <i>Nano Porous Hollow Fiber Regenerative Chemical Filter -</i>	0.989	0.000
Congressional Add: <i>Chemical and Biological Resistant Clothing</i>	0.000	1.593
Congressional Add: <i>Botulinum Toxin Treatment Therapy</i>	0.000	0.797
Congressional Add: <i>Contaminated Human Remains Pouch</i>	0.000	1.593
Congressional Add: <i>PaintShield for Protecting People from Microbial Threats</i>	0.000	1.992
Congressional Add Subtotals for Project: CI2	42.714	16.630
Congressional Add Totals for all Projects	42.714	16.630

Change Summary Explanation

Funding: N/A - Adjustments less than 10% of total program.

Schedule: N/A

Technical: N/A

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program								DATE: February 2010			
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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
CB2: <i>CHEMICAL BIOLOGICAL DEFENSE (APPLIED RESEARCH)</i>	102.599	110.955	88.897	0.000	88.897	100.243	97.979	90.686	91.554	Continuing	Continuing

A. Mission Description and Budget Item Justification

This project (CB2) provides physical applied research to develop future, multi-disciplinary, multi-functional capabilities in life sciences, physical sciences, environmental sciences, mathematics, cognitive sciences, and engineering. Efforts in this project support the seamless integration of state-of-the-art-technologies into a collection of systems across the spectrum of capabilities required to support chemical and biological defense missions, including specific research to develop defensive capabilities against non-traditional agents (NTAs). Starting in FY11, all NTA-dedicated research will be re-aligned into specific capability areas within this project in order to ensure a focused effort on this high priority area. Capability areas in this project include: detection; detection for NTAs; information systems technology; protection/hazard mitigation; protection/hazard mitigation for NTAs; threat agent science; and threat agent science for NTAs. Detection focuses on developing technologies for standoff and point detection and identification of chemical and biological agents. Information systems technology focuses on advanced warning and reporting, hazard prediction and assessment, simulation analysis and planning, and systems performance modeling. Protection and hazard mitigation focuses on providing technologies that protect and reduce the chemical/biological threat or hazard to the Warfighter, weapons platforms, and structures. Threat agent science is devoted to characterizing threat agents and the hazards they present in terms of agent fate in the environment, toxicology, pathogenicity and the development of simulants, especially with regard to NTAs. This project focuses on horizontal integration of CB defensive technologies in support of the Joint Services.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.	0.000	1.467	0.000	0.000	0.000
2) Protection	5.716	0.000	0.000	0.000	0.000

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 2: <i>Applied Research</i>	R-1 ITEM NOMENCLATURE PE 0602384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (APPLIED RESEARCH)</i>	PROJECT CB2: <i>CHEMICAL BIOLOGICAL DEFENSE (APPLIED RESEARCH)</i>				
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
of system alternatives and initiate efforts addressing specific technological gaps for COLPRO development. <i>FY 2010 Plans:</i> This effort re-aligned to Protection and Hazard Mitigation.						
8) Protection & Hazard Mitigation Innovative Systems Concepts and Analysis: Development and systems analysis of novel system concepts for chemical and biological protection of occupants of buildings and platforms that integrates emerging technologies. <i>FY 2010 Plans:</i> Investigate alternate system solutions and technologies for Collective Protection (COLPRO). Technologies include micro fine detoxifying aerosol fogs to facilitate entry and mitigate cross contamination into the COLPRO system, internal self-detoxifying surfaces for walls and ductwork, expedient retrofit kits, self-detoxifying and expedient strippable coatings, rapid isolation and purge schemes, and novel and innovative air flow and re-circulation schemes.		0.000	1.131	0.000	0.000	0.000
9) Protection & Hazard Mitigation Lightweight Integrated Fabric: Development of lightweight chemical and biological protective textiles that can be used as an integrated combat duty uniform. <i>FY 2010 Plans:</i> Support assessment of integrated fabric concurrent with the Individual Protection Advanced Technology Demonstration (IP Demo - see Budget Activity 3, Project TT3, Experiment and Technology Demonstrations), which will support the Lightweight CB Ensemble (LCBE), and incorporate lessons into further development of integrated fabric. Continue work on fabric residual life indicators and agent indicators that can be network enabled. Continue development of polymer membranes with		0.000	6.614	1.546	0.000	1.546

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p>Human Performance Prediction and Assessment: Analysis and modeling of human performance in chemical and biological protective ensembles in order to determine design priorities and trade-offs.</p> <p><i>FY 2010 Plans:</i> Support assessment of integrated fabric concurrent with the Individual Protection Advanced Technology Demonstration, which will support the Uniform Integrated Protective Ensemble (UIPE), and incorporate lessons into further development of human performance prediction and assessment. Continue refining human performance parameters for various Warfighter subgroups in the performance of their mission when CB protective systems are employed. Continue work to develop an overall comfort and performance model for CB protective equipment. Initiate anthropometric sizing study to support size tariff development.</p> <p><i>FY 2011 Base Plans:</i> Incorporate lessons learned from the Individual Protection Advanced Technology Demonstration, which will support the Uniform Integrated Protective Ensemble (UIPE), and incorporate lessons into further development of human performance prediction and assessment. Complete human performance model for CB protective equipment. As a result of the IP Demo, transition model data and analysis to individual protection advanced development programs. Continue anthropometric sizing study to support size tariff development.</p>						
<p>12) Protection & Hazard Mitigation</p> <p>Low-Burden Air Purifying Respirator: Development and analysis of design alternatives for chemical and biological air-purifying respirators to provide enhanced protection with lower physiological burden and improved interface with mission equipment.</p> <p><i>FY 2010 Plans:</i> Support assessment of integrated fabric concurrent with the Individual Protection Advanced Technology Demonstration, which will support the Uniform Integrated Protective Ensemble (UIPE),</p>		0.000	1.976	2.590	0.000	2.590

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p>novel, low pressure drop, HEPA filter, which provides increased dust capacity and extended filter life through the use of irregularly shaped high surface area submicron fibers.</p> <p><i>FY 2011 Base Plans:</i> Continue development of reactive membrane and regenerative post treatment media technologies for applications in building protection and vehicular/platform systems for Major Defense Acquisition Programs (MDAP).</p>						
<p>14) Protection & Hazard Mitigation</p> <p>General Purpose Formulations for Decontamination: Development and improvement of chemical and biological decontamination formulations that are compatible with the current family of decontamination systems.</p> <p><i>FY 2010 Plans:</i> Continue solid oxidant and green surfactant efforts resulting from alternative process research that emphasize dual-use technologies. Initiate focused enzymatic decontamination approaches.</p> <p><i>FY 2011 Base Plans:</i> Complete development, testing and transition of solid oxidant and green surfactant to support advanced development programs such as the Hazard Mitigation for Material and Equipment Restoration (HaMMER) Advanced Technology Demonstration (see Budget Activity 3, Project TT3, Experiment & Technology Demonstrations), also known as the Decontamination Family of Systems Demonstration. Continue focused enzymatic decontamination development.</p>		0.000	1.866	2.830	0.000	2.830
<p>15) Protection & Hazard Mitigation</p> <p>Decontamination System-of-Systems: Development and analysis of non-traditional decontamination technologies and approaches which gain significantly improved effectiveness by complementary application.</p>		0.000	2.553	4.348	0.000	4.348

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2010 Plans:</i> Complete development of self-detoxifying coatings, agent disclosure spray efforts, and strippable coating efforts and transition products in advanced development programs such as the Hazard Mitigation for Material and Equipment Restoration (HaMMER) Advanced Technology Demonstration. Continue investigation of microwave interaction with coating embedded particles and functionalities for directed energy decontamination. Complete work on functionalized photocatalytic materials. Initiate formulation development of a Decontamination Family of Systems that allow optimized formulation adjustment at point-of-use.</p> <p><i>FY 2011 Base Plans:</i> Develop data to define performance envelop of system components and transition to HaMMER. Initiate a study on impact of application methods of decontaminants to complex surfaces.</p>						
<p>16) Protection & Hazard Mitigation</p> <p>Smart Hazard Mitigation: Development of decontamination technologies that sense, respond (decontaminate) and signal in the presence of chemical and biological contamination.</p> <p><i>FY 2010 Plans:</i> Complete feasibility studies on the use of surface-modified nanoporous beads as encapsulation delivery devices for decontaminants. Continue development of molecular switches that respond and react to the presence of CB agents and signal results. Initiate development of rotaxane chemistry as artificial tunable G and V receptors that sense and react to chemical agents.</p> <p><i>FY 2011 Base Plans:</i> Continue development of molecular switches that respond and react to the presence of CB agents and signal results. Continue development of rotaxane chemistry as artificial tunable G and V receptors that sense and react to chemical and biological agents.</p>		0.000	1.787	1.388	0.000	1.388

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p>low volatility Chemical Warfare Agents (CWAs). Completed development of toxicokinetic and toxicodynamic models initiated in FY08.</p> <p><i>FY 2010 Plans:</i> Refine and standardize exposure and analytical methods for evaluation of percutaneous exposure to selected low volatility CWAs and high priority NTAs. Assess established contact and inhalation hazard methodologies for applicability to next-generation chemical warfare agents and refine as evaluation indicates. Set milestones and begin research on hazard assessment for more chemical agents. Complete development of exposure and analytic methods for selected very low volatile chemical threat agents. Complete studies and publish report on human health risk assessment exposure standard for medical applications associated with contact hazards of low volatility CWAs. Expand previous toxicokinetic and toxicodynamic efforts on a representative spore-forming Biological Weapons Agents (BWA) to include other BWAs, both spore-forming and non spore-forming. Assess the validity of expanding the viral agents model. Investigate human toxicity operational contact hazard assessment, and the effects of alternate toxicological pathways on the overall physiological impacts of high priority NTAs.</p> <p><i>FY 2011 Base Plans:</i> Continue research efforts on BWA toxicokinetic and toxicodynamic modeling. All NTA-related efforts re-aligned to Threat Agent Science NTA within this Budget Activity.</p>						
22) Threat Agent Science		4.990	8.847	0.079	0.000	0.079
Agent Fate: Characterizes fate of chemical and biological material on operationally relevant surfaces; information obtained from the study of particular agents will be used in core programs to assist detection, information systems, and protection and hazard mitigation activities.						

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B. Accomplishments/Planned Program (\$ in Millions)								
				FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
surfaces, including the impact of ambient conditions (e.g., temperature, relative humidity) and mechanical disturbances. All NTA-related efforts re-aligned to Threat Agent Science NTA within this Budget Activity.								
23) Threat Agent Science Accelerating Agent Sciences: Accelerates CB defense research and development by coupling computational methods and experimental approaches. <i>FY 2009 Accomplishments:</i> Continued CWA Quantum-Chemical Modeling (QCM) simulant design and selection methodology; simulant design and selection methodology efforts will be re-aligned to Agent Characterization and Simulant Development in FY10. Completed QCM dataset implementation to establish Quantitative Structure Activity Relationship (QSAR) between NTAs and surfaces/materials of operational interest. Utilized expertise and baseline against well-characterized substrates and move toward human toxicology QSAR toolsets. Integrated computational chemistry capabilities into experimental planning and data utilization work. <i>FY 2010 Plans:</i> Integrate research in computational techniques with existing computational toxicology, such as, shape signatures, and existing molecular dynamics capabilities to enhance agent fate, physiological response, simulant experiments and predictive modeling. Initiate work providing near term benefits, such as, computational toxicology. Complete CWA QCM development and maturation capability baseline for CWA interactions. Apply Quantum Chemical Modeling to develop and accelerate computationally obtained datasets and QSARS derived from the QCM data to highest priority NTA interactions and toxicology. <i>FY 2011 Base Plans:</i> All NTA-related efforts re-aligned to Threat Agent Science NTA within this Budget Activity.				4.482	3.861	0.000	0.000	0.000

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2011 Base Plans:</i> Continue BWA research to improve understanding of the relationship of genotype variations on organism virulence, infectivity, and persistence. Sustain efforts to support T&E applications by continued development of CWA and BWA simulants and refine simulant application by expanding agent-simulant correlation studies. All NTA-related efforts re-aligned to Threat Agent Science NTA within this Budget Activity.</p>						
<p>25) Threat Agent Science NTA</p> <p>Threat Agent Science NTA: Provides enabling science and technology which informs development and testing of NTA defense technology such as detection, decontamination, protection, hazard assessment, and more.</p> <p><i>FY 2011 Base Plans:</i> Establish human NTA operational toxicity estimates and interim human health risk assessments. Characterize the effects of alternate toxicological pathways. Expand agent fate studies to additional agent-substrate interactions. Correlate agent adsorption/absorption coefficients to chemical properties. Expand research on NTA liquid and solid phase transport to include re-suspension of particulates. Apply computational tools to identify data requirements and accelerate QSAR application to NTA interactions with operational substrates and toxicology issues. Correlate human effects to contact with operationally-relevant surfaces. Further research on NTA chemistry. Continue development of NTA simulants and simulant correlation studies.</p>		0.000	0.000	17.200	0.000	17.200
<p>26) Information Systems Technology</p> <p>Sensor Data Fusion: Emphasis on developing scientific techniques for fusing disparate information from multiple sources for insertion into the Joint Effects Model (JEM), Joint Warning and Reporting Network (JWARN), and Joint Operational Effects Federation (JOEF).</p>		4.980	0.000	0.000	0.000	0.000

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2009 Accomplishments:</i> Completed testing and validation and verification (V&V) of first-generation outdoor Source Term Estimation (STE), Hazard Refinement (HR) and Sensor Placement Tool (SPT) algorithms. Completed development, testing and V&V of building interior STE and HR algorithms. Initiated development of advanced STE, HR and SPT tools for use in complex environments (e.g., variable terrain, urban, water.) Completed biological background model development to reduce sensor false alarms and incorporate a first generation model into virtual environment software. Initiated development of a tool that continuously refines and updates the contamination footprint through rapid assimilation of limited and disparate information into meteorological, transport and dispersion, and virtual environment models.</p> <p><i>FY 2010 Plans:</i> Sensor Data Fusion efforts re-aligned to Advanced Warning and Reporting.</p>						
<p>27) Information Systems Technology</p> <p>Battle Space Management: Emphasis on development of collaborative information management technologies for insertion into the Joint Warning and Reporting Network (JWARN) and Joint Operational Effects Federation (JOEF) acquisition programs.</p> <p><i>FY 2009 Accomplishments:</i> Integrated Sensor Data Fusion (SDF) and source term location technologies into JEM and JOEF programs. Investigated and began development of next generation technologies and net-centric enterprise integration capabilities. Explored nano, bio, information technology and cognitive science solutions.</p> <p><i>FY 2010 Plans:</i> Battle Space Management efforts re-aligned to Advanced Warning and Reporting.</p>		2.990	0.000	0.000	0.000	0.000

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B. Accomplishments/Planned Program (\$ in Millions)								
				FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
programs. Finalize development of a tool that continuously refines and updates the contamination footprint through rapid assimilation of limited and disparate information into meteorological, transport and dispersion, and virtual environment models. Initiate development of route planning and evacuation/shelter-in-place decision aids.								
29) Information Systems Technology Hazard Prediction and Assessment: Improve battlespace awareness by accurately predicting hazardous material releases, atmospheric transport and dispersion, and resulting human effects. Develop predictive capability for the source term of releases of CB and industrial materials to include counterproliferation, CB weapons, accidents and ground effects from ballistic missiles. <i>FY 2009 Accomplishments:</i> Expanded and improved data assimilation techniques to develop a multi-scale, four-dimensional model. Continued development of advanced numerical weather prediction capabilities. Initiated optimization of methods to significantly improve performance of transport and dispersion hazard models for JEM. Developed advanced modeling capability for chemical, biological, and industrial source models (IFAC, ITRANS, and CBFAC). <i>FY 2010 Plans:</i> Initiate development of a missile intercept module for integration with JEM. Continue optimization of methods to significantly improve performance of transport and dispersion hazard models for JEM in both open air and urban environments using Second Order Closure Puff Atmospheric Transport and Dispersion (SCIPUFF AT&D) and Micro-Stationary Wind Fit with Turbulence (Micro-SWIFT). Continue advancing modeling techniques for chemical, biological, and industrial source models IFAC, ITRANS, and CBFAC. Continue experimental verification of models by way of small scale tests initiated in FY09.				2.257	4.942	3.030	0.000	3.030

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p>area in FY10. Closed out decision support data inscription technology to support change in advanced development program office priority. Continued distributed modeling research.</p> <p><i>FY 2010 Plans:</i> CBDP Decision Capability efforts re-aligned to Simulation Analysis and Planning.</p>						
<p>31) Information Systems Technology</p> <p>Chemical and Biological Warfare Effects on Operations: Develop the science behind the modeling and simulation of operations at the strategic, operational and tactical level in a CBRN environment for mobile forces, tactical aircraft, naval operations and fixed sites.</p> <p><i>FY 2009 Accomplishments:</i> Delivered methodology for CB effects on mobile and shipboard forces models to JOEF. Refined design and expanded prototype system for consequence management and incident management inclusions in consequence systems. Refined and expanded methodology for CBRN decision support tools.</p> <p><i>FY 2010 Plans:</i> Chemical and Biological Warfare Effects on Operations efforts re-aligned to Simulation Analysis and Planning.</p>		3.986	0.000	0.000	0.000	0.000
<p>32) Information Systems Technology</p> <p>Simulation Analysis and Planning: Develop decision support tools and information management capabilities for planning and real-time analysis to determine and assess operational effects, risks, and impacts of CBRN incidents on decision making. Focus areas include consequence management, human knowledge management, health/human effects modeling including casualty estimation, and fusion of diseases surveillance data.</p>		0.000	6.300	7.395	0.000	7.395

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2010 Plans:</i> Refine and update secondary infection models and NBC Casualty Resource Estimation Support Tool (NBC CREST) human effects models to reflect revision of NATO's Allied Medical Publication 8 (AMedP-8). Initiate development of casualty estimation methodology for CBRN agents including Non-Traditional Agents. Develop methodologies to improve the calculation of medical countermeasures effects in casualty estimation models. Improve CBRN medical resource planning tools. Continue development of contagious and infectious disease models. Continue development of particle size distribution health effects based on basic and applied threat agent science research efforts. Continue development and improvement of methodologies to apply CB operational effects in tactical, operational and strategic level models for mobile forces, shipboard modeling, fixed sites and tactical aircraft. Continue development of Incident Management/Consequence Management (IM/CM) tools and capabilities. Initiate studies to identify and investigate existing syndromic/disease surveillance systems and early detection capabilities. Continue validation and verification (V&V) effort for medical modeling efforts aimed at transitioning to advanced development efforts. Continue refinement and expansion of decision support tools for advanced development efforts. Complete distributed modeling research.</p> <p><i>FY 2011 Base Plans:</i> Complete development of refined versions of secondary infection models and human effects models to reflect revision of NATO's AMedP-8. Initiate development of additional casualty estimation modules for agents not in NATO's AMedP-8, including Non-Traditional Agents. Continue development of contagious/infectious disease models. Continue developing efforts aimed at integrating CB operational effects in tactical and operational level models for mobile forces, shipboard modeling, fixed sites and tactical aircraft. Further develop IM/CM tools and capabilities. Initiate development of capabilities that leverage and integrate existing early detection and disease surveillance data for inclusion into advanced development efforts.</p>						
33) Information Systems Technology		0.000	3.073	3.502	0.000	3.502

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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2011 Base Plans:</i> Complete algorithm development to increase range capabilities and reduce false positives. Complete work on first generation active infrared (IR) standoff biological classification capabilities. Complete evaluation and assessment of technology for scattering optical techniques, non-scattering optical standoff techniques, and off-gassing for down-selection of breadboard design. All NTA-related efforts re-aligned to Detection NTA within this Budget Activity.</p>					
<p>36) Detection NTA Primary focus is to assess the potential of optical technologies to meet the needs to detect the presence of NTAs. <i>FY 2011 Base Plans:</i> Complete a scientific analysis on the technical impacts of the presence of agents on surfaces due to the presence of NTAs. Complete assessment of chemical fate of chemicals in potable water. Continue feasibility development of plant sentinel concept. Complete design of first generation chemical standoff detection and identification capabilities. Initiate development from technology models to meet the needs to detect contamination on surfaces in a post decontamination application.</p>	0.000	0.000	12.000	0.000	12.000
Accomplishments/Planned Programs Subtotals	102.599	110.955	88.897	0.000	88.897

C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011 Base</u>	<u>FY 2011 OCO</u>	<u>FY 2011 Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To Complete</u>	<u>Total Cost</u>
• CB1: <i>CHEMICAL/BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	23.871	35.475	31.041		31.041	32.670	36.744	37.688	38.458	Continuing	Continuing
• CB3: <i>CHEMICAL BIOLOGICAL DEFENSE (ATD)</i>	19.567	25.297	15.410		15.410	21.450	26.120	36.775	37.148	Continuing	Continuing
	25.761	13.307	11.875		11.875	11.267	11.160	0.000	0.000	Continuing	Continuing

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C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011 Base</u>	<u>FY 2011 OCO</u>	<u>FY 2011 Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To Complete</u>	<u>Total Cost</u>
• TE3: <i>TEST & EVALUATION (ATD)</i>											
• TT3: <i>TECHBASE TECHNOLOGY TRANSITION</i>	8.127	7.357	4.504		4.504	8.117	8.169	8.390	8.528	Continuing	Continuing

D. Acquisition Strategy

N/A

E. Performance Metrics

N/A

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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
CI2: <i>CONGRESSIONAL INTEREST ITEMS (APPLIED RESEARCH)</i>	42.714	16.630	0.000	0.000	0.000	0.000	0.000	0.000	0.000	Continuing	Continuing

A. Mission Description and Budget Item Justification

The efforts this project include congressional interest programs for FY09 and FY10.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010
Congressional Add: Rapid Forensic Evaluation of Microbes in Biodefense <i>FY 2009 Accomplishments:</i> Continued research program to develop an ultra-sensitive single application detection method that can be used for a range of Bioterrorism agents.	0.989	0.000
Congressional Add: Chem/Bio IR Detection System <i>FY 2009 Accomplishments:</i> Continued research to investigate an electric-field focusing approach, combined with optically transparent filters, to be used for spore capture and identification. <i>FY 2010 Plans:</i> Develop an advanced chemical and biological detection system using a common platform to include detection of emerging novel agents and toxic industrial chemicals	1.186	1.892
Congressional Add: Zumwalt National Program for Countermeasures to Bio Chem Threats	1.187	0.000

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B. Accomplishments/Planned Program (\$ in Millions)		
	FY 2009	FY 2010
<i>FY 2009 Accomplishments:</i> Continued research to improve model development related to atmospheric sciences and environmental modeling.		
Congressional Add: HyperAcute Vaccine Development <i>FY 2009 Accomplishments:</i> Continued research by testing vaccine efficacy in a mouse model for correlates of immunity and protection from live virus challenge. <i>FY 2010 Plans:</i> Continuation of research from FY09.	2.373	3.585
Congressional Add: Antibody-based Therapeutic against Smallpox <i>FY 2009 Accomplishments:</i> Continued testing with the goal of generating a combinational therapeutic of human mAbs to several neutralizing VACV proteins, that confer the highest degree of protection against vaccinia, smallpox, monkeypox, and other orthopoxvirus infections.	0.791	0.000
Congressional Add: Novel Viral Biowarfare Agent Identification and Treatment (NOVBAIT) <i>FY 2009 Accomplishments:</i> Continued research to find small molecules that inhibit the assembly of capsids by viruses of high biowarfare potential, thereby inhibiting their replication and neutralizing infection.	3.955	0.000
Congressional Add: Mixed Oxidants for Chemical and Biological Decontamination	2.769	0.000

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B. Accomplishments/Planned Program (\$ in Millions)		
	FY 2009	FY 2010
<i>FY 2009 Accomplishments:</i> Continued research begun in FY08.		
Congressional Add: Bio Surety Development and Management Program - <i>FY 2009 Accomplishments:</i> Continued the research and analysis from FY08.	1.186	0.000
Congressional Add: Countermeasures to Chemical/Biological Control-Rapid Response - <i>FY 2009 Accomplishments:</i> Continued research from FY08.	2.372	0.000
Congressional Add: Multiple Applications for Light Activated, Reactive Materials for Protection of Warfighter, First Responder, and Public Health - <i>FY 2009 Accomplishments:</i> Developed protective applications for first responders of all types.	1.582	0.000
Congressional Add: Chemical Biological Preparedness Center for Advanced Development of Mobile Rapid Response Prototype - <i>FY 2009 Accomplishments:</i> Developed a mobile, forward deployable, medical capacity that would respond to bio-terrorist incidents and other mass casualty incidents resulting from WMD, natural and technological disasters.	3.955	0.000
Congressional Add: Novel System for Developing Therapeutics against Botulism -	3.955	0.000

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B. Accomplishments/Planned Program (\$ in Millions)		
	FY 2009	FY 2010
<i>FY 2009 Accomplishments:</i> Conducted research to discover new therapeutics against Botulism.		
Congressional Add: Ultra-Rapid Next Generation Pathogen Identification - <i>FY 2009 Accomplishments:</i> Developed pathogen identification capabilities.	1.978	0.000
Congressional Add: Preventing Long-Term Brain and Lung Damage Caused by Battlefield Trauma Project - <i>FY 2009 Accomplishments:</i> Conducted research to determine new techniques to prevent brain and lung damage.	2.868	0.000
Congressional Add: Chemical Agent Fate Appropriate Response Tool - <i>FY 2009 Accomplishments:</i> Conducted research to create a systematic approach for to the development of a comprehensive operational agent fate model/tool that provides recommendations on the appropriate response to contamination events. <i>FY 2010 Plans:</i> Continue research from FY09.	1.582	1.593
Congressional Add: Multivalent Marburg/Ebola Vaccine - <i>FY 2009 Accomplishments:</i> Conducted research in the development of a multivalent Marburg/Ebola vaccine.	3.461	0.000

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B. Accomplishments/Planned Program (\$ in Millions)		
	FY 2009	FY 2010
Congressional Add: Botulinum Neurotoxin Research - <i>FY 2009 Accomplishments:</i> Conducted research in the development of a new assay which is designed to detect Botulinum (A-G) in the environment and on exposed animals, humans, and culture cells. <i>FY 2010 Plans:</i> Continue research from FY09.	1.582	1.992
Congressional Add: Miniaturized Chemical Detector for Chemical Warfare Protection (ChemPen) - <i>FY 2009 Accomplishments:</i> Developed a ready for production MEMs FTIR absorption spectrometer to detect in seconds a wide range of nerve agents/TICs. <i>FY 2010 Plans:</i> Continuation of research from FY09.	1.581	1.593
Congressional Add: Continued Expansion of Prototypes for Destruction of Airborne Pathogen - <i>FY 2009 Accomplishments:</i> Continued development of methodologies for the destruction of aerosolized agents.	0.791	0.000
Congressional Add: Mismatch Repair Derived Antibody to Treat Staph Derived Bioweapon - <i>FY 2009 Accomplishments:</i> Continued research begun in FY07 to develop fully human anti-Staphylococcus enterotoxin B (SEB) monoclonal antibodies (mAbs) that can neutralize >1000 times the human LD50 of the toxin.	1.582	0.000

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B. Accomplishments/Planned Program (\$ in Millions)		
	FY 2009	FY 2010
Congressional Add: Nano Porous Hollow Fiber Regenerative Chemical Filter - <i>FY 2009 Accomplishments:</i> Conducted research in the application of nanotechnology to chemical filter design.	0.989	0.000
Congressional Add: Chemical and Biological Resistant Clothing <i>FY 2010 Plans:</i> Develop a material capable of simultaneously being lightweight, robust, breathable, and resistant to chemical and biological agents.	0.000	1.593
Congressional Add: Botulinum Toxin Treatment Therapy <i>FY 2010 Plans:</i> Develop new therapies for botulinum toxin poisoning and other bioterrorism threats.	0.000	0.797
Congressional Add: Contaminated Human Remains Pouch <i>FY 2010 Plans:</i> Conduct development activities for a contaminated human remains transportable container.	0.000	1.593
Congressional Add: PaintShield for Protecting People from Microbial Threats <i>FY 2010 Plans:</i> Develop a paint coating technology, a cost-effective, interior paint platform that will render microbiological threats harmless upon contact.	0.000	1.992
Congressional Adds Subtotals	42.714	16.630

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C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011</u> <u>Base</u>	<u>FY 2011</u> <u>OCO</u>	<u>FY 2011</u> <u>Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To</u> <u>Complete</u>	<u>Total Cost</u>
• CI1: <i>CONGRESSIONAL INTEREST ITEMS (BASIC RESEARCH)</i>	8.090	20.036	0.000		0.000	0.000	0.000	0.000	0.000	Continuing	Continuing
• CI3: <i>CONGRESSIONAL INTEREST ITEMS (ATD)</i>	46.971	18.622	0.000		0.000	0.000	0.000	0.000	0.000	Continuing	Continuing

D. Acquisition Strategy

N/A

E. Performance Metrics

N/A

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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
TB2: <i>MEDICAL BIOLOGICAL DEFENSE (APPLIED RESEARCH)</i>	50.485	53.930	43.858	0.000	43.858	50.866	51.077	51.051	51.959	Continuing	Continuing

A. Mission Description and Budget Item Justification

This project (TB2) funds applied research of vaccines, therapeutic drugs, and diagnostic capabilities to provide effective medical defense against validated biological threat agents including bacteria, toxins, and viruses. Innovative biotechnology approaches will be incorporated to advance medical systems designed to rapidly identify, diagnose, prevent, and treat disease due to exposure to biological threat agents will be advanced. Categories of this project include core science efforts in biological defense capability areas, such as Pretreatments, Diagnostics, and Therapeutics. Medical Science and Technology (S&T) efforts in this Budget Activity refine promising medical initiatives identified in Budget Activity 1, resulting in the development of countermeasures to protect against and treat the effects of exposure to chemical, biological, and radiological (CBR) agents.

This project also includes efforts such as the Transformational Medical Technologies Initiative (TMTI). The Transformational Medical Technologies Initiative (TMTI) was launched to respond to the threat of emerging or intentionally bioengineered biological threats. TMTI's mission is to protect the Warfighter from genetically engineered biological threats by providing a rapid response capability from identification of pathogens to the delivery of medical countermeasures. This mission is accomplished through two main efforts: 1) developing broad spectrum (multi-agent) therapeutics against biological warfare (BW) agents (e.g. one drug that treats multiple agents); and 2) developing platform technologies to assist in the rapid development of medical countermeasures (MCMs) in response to BW agents (e.g. developing new and innovative ways to mass produce drugs in the event of a biological incident).

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.	0.000	0.733	0.000	0.000	0.000
2) Diagnostics	6.594	7.197	6.994	0.000	6.994

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2010 Plans:</i> This effort re-aligned to Vaccine Platforms and Research Tools.						
4) Pretreatments Multiagent Vaccine Platforms: Construct multi-agent vaccine platforms and formulations capable of expressing multiple protein antigens from multiple pathogens, and evaluate in animal models. <i>FY 2009 Accomplishments:</i> Further assessed candidate multi-agent vaccines in animal models, and considered the inclusion of alternative agents. Explored novel platforms and vaccine formulations. Evaluated effectiveness in animal models. <i>FY 2010 Plans:</i> Effort re-aligned to Vaccine Platforms and Research Tools.		1.364	0.000	0.000	0.000	0.000
5) Pretreatments Vaccine Research Support: Identify the elements of a vaccine formulation that are necessary for an effective host immune response that confers protection against biothreat agents. <i>FY 2009 Accomplishments:</i> Further characterized immune correlates of protection elicited by alphavirus (WEE/VEE/EEE) and filovirus vaccines in animal models. Optimized alphavirus and filovirus antibody-based assays and evaluated their ability to predict protection. Explored additional intracellular pathogen antigens using animal model systems including the use of alternative vaccine delivery platforms for protection. Further evaluated the protective efficacy of BoNT components in small animal models. Extended the characterization of non-protective antigen vaccine candidates to additional small animal models. Pursued the use of immune stimulating protein fragments (peptides) or immune cell targeting peptides to enhance vaccine efficacy in animal models.		6.463	0.000	0.000	0.000	0.000

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2010 Plans:</i> Efforts re-aligned to Viral Vaccines and Bacterial/Toxin Vaccines.						
6) Pretreatments Bacterial/Toxins Vaccines: Generate novel or improved vaccines against bacterial and toxin biothreat agents, and demonstrate preliminary efficacy in small animal models. Identify correlates of protective immunity in animals models. <i>FY 2010 Plans:</i> Test the efficacy of Burkholderia vaccine candidates against aerosol challenge in small animal models. Begin to determine the therapeutic regimen needed in conjunction with a vaccine to eliminate residual Burkholderia organisms and begin evaluation of the immune response elicited by the vaccine. Use comparative animal studies to test the efficacy of disease inactivated, but metabolically active vaccine candidates against Brucella species. Begin to compare the ability of the disease inactivated, but metabolically active vaccine candidates to protect mice against aerosol challenge with distinct strains of Brucella following oral immunization. Continue to test the immune stimulation and effectiveness of novel anthrax vaccines (e.g., multi-component genetically altered vaccines composed of spore antigens, etc.) to combat emerging and genetically engineered strains. Initiate studies aimed at generating a second-generation vaccine that protects against aerosolized Type A Francisella tularensis. <i>FY 2011 Base Plans:</i> Continue aerosol efficacy studies in mice for Brucella and Burkholderia vaccine candidates. Work to improve the efficacy of the most promising vaccine candidates against Burkholderia and Brucella by initiating studies that vary the route of immunization, dose and vaccination schedule. Begin investigating whether the efficacy of the Brucella and Burkholderia vaccine candidates can be approved by co-administering the vaccines with nonspecific stimulators of the immune response (i.e., adjuvants). Test the ability of antibiotics to remove residual Burkholderia from vaccinated animals to		0.000	2.948	5.254	0.000	5.254

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
the MIMIC in response to biodefense vaccines, and develop a sensitive fluorescent-based assay to assess the functionality of the antibodies generated. Adapt the MIMIC to function as an infectious disease model for alphaviruses and filoviruses. Use these MIMIC in infectious disease models to begin to define human correlates of protective immunity against alphaviruses and filoviruses. Initiate studies to develop methodologies that render different types of vaccine platforms (i.e., viral vector, inactivated virus, virus like particles, and attenuated bacteria, etc.) stable in variable and extreme temperatures.						
9) Therapeutics Therapy for Ebola and Marburg Virus Infections: Identify, optimize and evaluate lead candidate therapeutics for efficacy against Filovirus infections, specifically Ebola and Marburg Viruses. <i>FY 2009 Accomplishments:</i> Completed proof-of-concept studies for lead candidate technologies.		0.811	0.000	0.000	0.000	0.000
10) Therapeutics Viral Therapeutics: Identify, optimize and evaluate lead candidate therapeutics for efficacy against viral pathogens. <i>FY 2009 Accomplishments:</i> Determined the ability of heavy metal nanoparticle-based therapeutics to inhibit viral infection in a laboratory model system. Conducted proof-of-concept studies aimed at identifying therapeutic candidates for poorly characterized threats. Continued supporting therapeutics effective against well characterized threat agents towards advanced development. Screened multiple compound libraries for small molecule inhibitors of designated viral pathogens.		0.430	2.067	1.600	0.000	1.600

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2010 Plans:</i> Initiate drug discovery for a second novel orthopox drug with a mechanism distinct from ST-246, a low-molecular-weight compound that is active against multiple orthopoxviruses. Expand drug discovery efforts for alphaviruses (VEE, EEE, and WEE). Establish clinical protocols to obtain human clinical samples from filovirus outbreaks in the Democratic Republic of the Congo. Test and evaluate lead candidate therapeutic compounds in relevant animal challenge models. Continue testing of heavy metal nanoparticle-based therapeutics for the ability to prevent viral infection in animal models. Identify lead compounds from small molecule library screening and optimize their action through medicinal chemistry. Test and evaluate small protein fragments to determine if their ability to prevent a virus from binding to cells represents a viable therapeutic interdiction point for designated viral pathogens.</p> <p><i>FY 2011 Base Plans:</i> Identify FDA-approved drug combinations with efficacy against alphavirus infection. Identify and develop small molecule inhibitors to specific host factors required for alphavirus pathogenesis. Conduct structure-based screening of chemical libraries to identify inhibitors of alphavirus proteins. Utilize medicinal chemistry to optimize antiviral activity of lead compounds. Identify therapeutic inhibitors of orthopoxvirus infection by targeting required host and viral tyrosine phosphatases.</p>						
<p>11) Therapeutics</p> <p>Bacterial Therapeutics: Identify, optimize and evaluate lead therapeutic candidates effective against designated bacterial threat agents.</p> <p><i>FY 2009 Accomplishments:</i> Completed initial evaluation of a single domain antibody that is smaller than conventional antibodies against plague, and extend the application to other related bacteria if successful. Screened small molecules that can prevent plague bacteria from injecting virulence factors into cells in the laboratory, and extend application of assay to other related bacteria. Balanced efforts to evaluate potential single</p>		5.418	4.110	4.100	0.000	4.100

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p>agent bacterial therapeutics with those having broad-spectrum activity. Identified and screened inhibitors of bacterial phosphatases for protective effects in cellular and animal models.</p> <p><i>FY 2010 Plans:</i> Complete evaluation of bacterial phosphatase inhibitors in a mouse model of plague infection. Test and evaluate lead candidate small molecules to determine their antimicrobial activity. Screen commercially available antimicrobial in advanced clinical development for their activity in the laboratory against bacterial threat agents.</p> <p><i>FY 2011 Base Plans:</i> Continue the identification of commercially available antimicrobials in advanced clinical development with laboratory assayed activity against bacterial threat agents. Assess compounds identified in high content imaging assays for their antimicrobial activity in relevant animal challenge models.</p>						
<p>12) Therapeutics</p> <p>Toxin Therapeutics: Identify, optimize and evaluate therapeutic candidates that are effective against biological toxin agents.</p> <p><i>FY 2009 Accomplishments:</i> Evaluated next generation monoclonal antibodies for laboratory and animal effectiveness against Botulinum Neurotoxin (BoNT). Characterized lead compounds for potency and specificity in laboratory models and animal models. Initiated development of inactive versions of BoNT substrates as therapeutics with the potential to restore nerve activity following neuromuscular paralysis. Developed a cell-based high-throughput screening system for BoNT therapeutics derived from mouse cells and embryonic stem cells. Evaluated immune-modifying compounds for pre- and post-exposure therapy for Staphylococcal Enterotoxin B (SEB) intoxication in laboratory and animal models.</p>		10.528	9.065	9.171	0.000	9.171

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2010 Plans:</i> Screen compound libraries utilizing a high-throughput screening system for BoNT therapeutics derived from mouse cells and embryonic stem cells. Test and evaluate lead candidate inhibitors in relevant laboratory and animal model systems of BoNT intoxication. Perform experimental analysis to clarify the contribution of protein modification of BoNT to its structure and biochemical activity as it relates to drug development. Conduct high-throughput screening of drug libraries to identify inhibitors of ricin toxicity.</p> <p><i>FY 2011 Base Plans:</i> Develop transgenic mice expressing genetically-encoded reporters of BoNT activity in neurons for use in high-throughput screening of BoNT therapeutics. Validate neurite outgrowth analysis for the identification of BoNT inhibitors. Identify host proteins responsible for BoNT light chain stabilization. Conduct co-crystallization studies of BoNT-inhibitor complexes. Perform experiments to determine toxicity and pharmacokinetics of selected ricin inhibitors. Identify host proteins involved in ricin dislocation as potential host-directed drug targets. Determine efficacy of identified ricin inhibitors in mice.</p>						
13) Transformational Medical Technologies Initiative		15.005	4.186	8.037	0.000	8.037
<p>Multiagent (Broad Spectrum) Medical Countermeasures (MCM): Builds upon basic research performed by existing performers and supports the efforts of new performers who are in the mid-drug discovery phase of drug development. Applied research efforts also include the investigation of existing drugs to explore their efficacy against BW agents. Assesses toxicity and efficacy in accordance with the product's intended use. Initiation of experiments to identify markers, correlates of protection, assays, and endpoints for further non-clinical and clinical studies. Develop a scalable and reproducible manufacturing process amenable to Food and Drug Administration (FDA) good manufacturing processes.</p>						

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2010 Plans:</i> Identified enabling and critical technologies, formulated appropriate technology plans and acquisition strategies, and determined their performance objectives. Initiated development of an information network to serve as the backbone for a rapid drug discovery and development capability. Supported development of platform technologies to higher levels of maturity. Genetic sequencing studies model the types and quantity of data needed for the identification of unknown pathogen ID, including a genomic survey for countermeasure targets and genetically engineering. Evaluated the information network to serve as the backbone for a rapid drug discovery and development capability. Pursued informatics to support analytical activities, event response, and science discovery. Initiated work on advanced manufacturing to enhance the rapid production of therapeutics.</p> <p><i>FY 2011 Base Plans:</i> Continue the development of host and pathogen based platforms to higher levels of maturity. Continue to explore pathogen identification and characterization capabilities, including genetic sequencing, integrate existing capabilities. Continue to assess future sequence and analysis needs to characterize advanced threats. Continue to integrate leading edge technologies with existing technologies to enhance pathogen characterization, target identification, countermeasure discovery and countermeasure evaluation platform areas.</p>					
Accomplishments/Planned Programs Subtotals	50.485	53.930	43.858	0.000	43.858

C. Other Program Funding Summary (\$ in Millions)

Line Item	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	FY 2012	FY 2013	FY 2014	FY 2015	Cost To Complete	Total Cost
• TB1: <i>MEDICAL BIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	15.086	16.782	14.352		14.352	15.499	14.845	14.402	14.672	Continuing	Continuing
	180.425	203.723	115.233		115.233	125.666	109.737	115.049	117.289	Continuing	Continuing

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program	DATE: February 2010
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APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 2: <i>Applied Research</i>	R-1 ITEM NOMENCLATURE PE 0602384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (APPLIED RESEARCH)</i>	PROJECT TB2: <i>MEDICAL BIOLOGICAL DEFENSE (APPLIED RESEARCH)</i>
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C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011</u> <u>Base</u>	<u>FY 2011</u> <u>OCO</u>	<u>FY 2011</u> <u>Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To</u> <u>Complete</u>	<u>Total Cost</u>
• TB3: <i>MEDICAL BIOLOGICAL DEFENSE (ATD)</i>											

D. Acquisition Strategy

N/A

E. Performance Metrics

N/A

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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
TC2: <i>MEDICAL CHEMICAL DEFENSE (APPLIED RESEARCH)</i>	35.008	40.418	33.648	0.000	33.648	36.327	36.500	37.475	38.150	Continuing	Continuing

A. Mission Description and Budget Item Justification

This project (TC2) funds applied research for the investigation of new medical countermeasures to include prophylaxes, pretreatments, antidotes, skin decontaminants and therapeutic drugs against identified and emerging chemical warfare threat agents to include a class of agents called Non Traditional Agents (NTAs). Starting in FY11, all NTA-dedicated research will be re-aligned into specific capability areas within this project in order to ensure a focused effort on this high priority area. Capability areas include: Pretreatments; pretreatments for NTAs; diagnostics; diagnostics for NTAs; therapeutics; and therapeutics for NTAs. Pretreatments includes researching prophylaxes to protect against chemical agents and NTAs. Diagnostics focuses on researching diagnostic tools that help identify exposure to chemical agents and NTAs. Therapeutics focuses on researching post-exposure countermeasures to protect against chemical agents and NTAs. Research and development efforts in this project focus on formulation and scale-up of candidate compounds.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.	0.000	0.507	0.000	0.000	0.000
2) Diagnostics Diagnostic Technologies: Focuses on developing state-of-the-art laboratory/fieldable methods that detect exposure to chemical warfare agents (CWA) (e.g., nerve agents and vesicants) in clinical samples. Identifies biomolecular targets that can be leveraged as analytical methodologies, as well as, laboratory and animal studies characterizing time-course and longevity of a particular analyte/ biomarker.	1.013	1.207	0.865	0.000	0.865

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2009 Accomplishments:</i> Completed alternative sample collection/extraction technologies, such as, solvent free extraction as part of a rapid screening method to verify exposure to CWAs. Evaluated the combined sample extraction and analysis procedure for pre- and post-CWA exposure to assess the feasibility of detecting chemical warfare analytes in hair samples from animals. Incorporated promising antibody diagnostics and molecular technologies for hand-held CWA diagnostic platforms developed under the Small Business Innovative Research (SBIR) program into the core program for further development.</p> <p><i>FY 2010 Plans:</i> Continue development of definitive diagnostic biomarkers for early detection of CWA exposure using several different analytical approaches. Develop pre-symptomatic diagnostic technologies for eventual incorporation into handheld devices in order to detect CWA exposures.</p> <p><i>FY 2011 Base Plans:</i> Continue to determine whether existing CWA biomarkers are appropriate for early detection and validation of CWA exposure in clinical samples. Determine if biomarkers that appear after exposure to sulfur mustard can be used to identify an appropriate treatment option prior to the onset of symptoms. Continue investigation of a novel surface plasmon resonance based sensor array and a phage library display to develop binding molecules as biomarkers of nerve agent exposure. All NTA-related efforts are re-aligned to Chemical Diagnostics NTA within this Budget Activity.</p>						
3) Chem Diagnostics NTA <i>FY 2011 Base Plans:</i> Continue studies to identify biomarkers to create an enhanced capability to pre-symptomatically diagnose NTA exposure. Continue method development for identification and validation of NTAs in clinical samples.		0.000	0.000	0.400	0.000	0.400
4) Pretreatments		10.685	9.883	5.980	0.000	5.980

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
for neuroprotection against nerve agent exposure. Develop therapeutic strategies to effectively minimize neurologic injuries resulting from exposure to CWAs by testing in silico and in vitro.						
9) Therapeutics Medical Toxicology (Non Traditional Agents (NTAs) and Other Agents): Investigates common mechanisms of agent injury. Determines the toxic effects of agents by probable routes of field exposure, as well as standard experimental routes. Physiological parameters and pathological assessment will be used to establish the general mode and mechanism(s) of toxicity. <i>FY 2009 Accomplishments:</i> Quantified the nature, scope, and time course of exposure/effects using biochemical, toxicological, physiological, and modeling methods as required for therapeutic and clinical strategy design. <i>FY 2010 Plans:</i> Investigate and study receptor effects of common and agent-specific mechanisms of NTA injury for therapeutic intervention. <i>FY 2011 Base Plans:</i> All NTA-related efforts are re-aligned to Chemical Therapeutics NTA within this Budget Activity.		1.783	2.756	0.000	0.000	0.000
10) Therapeutics Therapeutics for Non Traditional Agents (NTAs): Develops, assesses, evaluates, and validates therapeutics for treatment resulting from exposure to NTAs. <i>FY 2009 Accomplishments:</i> Evaluated pre-existing and new commercially-available compounds for respiratory and neurological injury in small animal models and began transition to large animal models (e.g. non-human primate).		8.818	13.156	0.000	0.000	0.000

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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p>Initiated testing of novel compounds as therapies in small animal models. Defined and optimized the utility of therapeutic agent-binding proteins against NTAs.</p> <p><i>FY 2010 Plans:</i> Further development and validation of animal models for testing clinical efficacy of therapeutics against NTAs. Identify binding characteristics of NTAs, as well as mitigate NTA toxicity by researching and developing novel therapeutics.</p> <p><i>FY 2011 Base Plans:</i> All NTA-related efforts are re-aligned to Chemical Therapeutics NTA within this Budget Activity.</p>					
<p>11) Chem Therapeutics NTA</p> <p>Develops, assesses, evaluates, and validates therapeutics for treatment resulting from exposure to NTAs.</p> <p><i>FY 2011 Base Plans:</i> Continue binding studies to support the design and synthesis of an improved reactivator. Continue evaluation of improved products to treat NTA exposure. Continue investigation of pathophysiological effects to identify debilitating syndromes caused by exposure to NTAs. Continue development of animal models for various routes of exposure to NTA. These models will be utilized to evaluate toxic effects of NTAs, behavioral changes, efficacy, and FDA animal rule approvals.</p>	0.000	0.000	13.000	0.000	13.000
Accomplishments/Planned Programs Subtotals	35.008	40.418	33.648	0.000	33.648

C. Other Program Funding Summary (\$ in Millions)

Line Item	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	FY 2012	FY 2013	FY 2014	FY 2015	Cost To Complete	Total Cost
• TC1: <i>MEDICAL CHEMICAL DEFENSE (BASIC RESEARCH)</i>	13.308	5.496	3.144		3.144	2.889	2.954	2.928	2.977	Continuing	Continuing

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011</u> <u>Base</u>	<u>FY 2011</u> <u>OCO</u>	<u>FY 2011</u> <u>Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To</u> <u>Complete</u>	<u>Total Cost</u>
• TC3: <i>MEDICAL CHEMICAL DEFENSE (ATD)</i>	21.641	28.971	29.134		29.134	30.401	30.546	31.356	31.877	Continuing	Continuing

D. Acquisition Strategy

N/A

E. Performance Metrics

N/A

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
TR2: <i>MEDICAL RADIOLOGICAL DEFENSE (APPLIED RESEARCH)</i>	0.525	2.897	2.884	0.000	2.884	1.904	2.855	1.913	1.903	Continuing	Continuing

A. Mission Description and Budget Item Justification

This project (TR2) funds applied research to develop medical countermeasures to protect the Warfighter against radiological exposure. Specifically, innovative technical approaches will be used to develop products to mitigate health consequences resulting from Acute Radiation Exposure (ARS) and Delayed Effects of Acute Radiation Exposure (DEARE). The research and development of medical countermeasures for radiation exposure will ultimately enhance the survivability of Warfighters and will serve to significantly minimize the development of acute radiation syndromes and subsequent health problems. Results of efforts funded under this project are collaboratively shared with other government agencies, while the Department of Defense maintains an emphasis on the development of pretreatments to protect military personnel who could be involved in responding to a radiological incident.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.	0.000	0.042	0.000	0.000	0.000
2) Radiological Medical Countermeasures Radiation Medical Countermeasures: Develop medical countermeasures to protect the Warfighter against radiological/nuclear exposure, to include developing both pretreatments (prophylaxis) and post-irradiation therapeutics against radiological/nuclear exposure. DoD is the only governmental agency currently developing medical prophylaxis to protect Warfighters and/or other responders in the event of a radiological incident.	0.525	2.855	2.884	0.000	2.884

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 2: <i>Applied Research</i>	R-1 ITEM NOMENCLATURE PE 0602384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (APPLIED RESEARCH)</i>	PROJECT TR2: <i>MEDICAL RADIOLOGICAL DEFENSE (APPLIED RESEARCH)</i>
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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2009 Accomplishments:</i> Down-select at least one promising drug candidate that has the ability to provide protection from the harmful effects of radiation exposure. Determine the pre-clinical efficacy of combined agents that confer protection or supportive medical care against the harmful effects of radiation exposure with minimal toxic side effects. Explore current Good Laboratory Practice (cGLP) test capability for selected candidate drugs against acute radiation syndrome (ARS) based on Food and Drug Administration's (FDA) animal testing requirements.</p> <p><i>FY 2010 Plans:</i> Evaluate mature and promising drug candidates for respiratory and gastrointestinal damage and repair, demonstrating efficacy, safety, and animal (rodents) survival exposed to lethal radiation for a future non-human primate (NHP) efficacy study. Identify common biochemical/physiological mechanisms for hematological, respiratory and gastrointestinal damage and repair, as well as, biology of cellular damage.</p> <p><i>FY 2011 Base Plans:</i> Continue to evaluate novel and FDA-approved drugs for efficacy against radiation exposure maintaining a focus on potential mechanisms of action. These studies will help identify biochemical/physiological mechanisms that could be exploited for expanding the scope of potential therapeutic approaches. Continue to focus approaches on the GI and lung injury related to radiation exposure. Continue evaluation and identification of unique, novel and promising biomarkers useful for biodosimetry and potential pathways for therapeutic approaches.</p>					
Accomplishments/Planned Programs Subtotals	0.525	2.897	2.884	0.000	2.884

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 2: <i>Applied Research</i>	R-1 ITEM NOMENCLATURE PE 0602384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (APPLIED RESEARCH)</i>	PROJECT TR2: <i>MEDICAL RADIOLOGICAL DEFENSE (APPLIED RESEARCH)</i>
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C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011</u> <u>Base</u>	<u>FY 2011</u> <u>OCO</u>	<u>FY 2011</u> <u>Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To</u> <u>Complete</u>	<u>Total Cost</u>
• TR1: <i>MEDICAL RADIOLOGICAL DEFENSE (BASIC RESEARCH)</i>	0.000	0.975	0.971		0.971	0.966	0.000	0.000	0.000	Continuing	Continuing
• TR3: <i>MEDICAL RADIOLOGICAL DEFENSE (ATD)</i>	4.859	2.403	0.957		0.957	0.966	1.922	2.901	2.927	Continuing	Continuing

D. Acquisition Strategy

N/A

E. Performance Metrics

N/A

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Exhibit R-2, RDT&E Budget Item Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 3: <i>Advanced Technology Development (ATD)</i>	R-1 ITEM NOMENCLATURE PE 0603384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ATD)</i>
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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
Total Program Element	307.351	299.680	177.113	0.000	177.113	197.867	187.654	194.471	197.769	Continuing	Continuing
CB3: <i>CHEMICAL BIOLOGICAL DEFENSE (ATD)</i>	19.567	25.297	15.410	0.000	15.410	21.450	26.120	36.775	37.148	Continuing	Continuing
CI3: <i>CONGRESSIONAL INTEREST ITEMS (ATD)</i>	46.971	18.622	0.000	0.000	0.000	0.000	0.000	0.000	0.000	Continuing	Continuing
TB3: <i>MEDICAL BIOLOGICAL DEFENSE (ATD)</i>	180.425	203.723	115.233	0.000	115.233	125.666	109.737	115.049	117.289	Continuing	Continuing
TC3: <i>MEDICAL CHEMICAL DEFENSE (ATD)</i>	21.641	28.971	29.134	0.000	29.134	30.401	30.546	31.356	31.877	Continuing	Continuing
TE3: <i>TEST & EVALUATION (ATD)</i>	25.761	13.307	11.875	0.000	11.875	11.267	11.160	0.000	0.000	Continuing	Continuing
TR3: <i>MEDICAL RADIOLOGICAL DEFENSE (ATD)</i>	4.859	2.403	0.957	0.000	0.957	0.966	1.922	2.901	2.927	Continuing	Continuing
TT3: <i>TECHBASE TECHNOLOGY TRANSITION</i>	8.127	7.357	4.504	0.000	4.504	8.117	8.169	8.390	8.528	Continuing	Continuing

A. Mission Description and Budget Item Justification

This program element (PE) demonstrates technologies that enhance the ability of U.S. forces to deter, defend against, and survive Chemical, Biological, and Radiological (CBR) warfare. This program element (PE) funds advanced technology development for Joint Service and Service-specific requirements in both medical and physical sciences CBR defense areas. The medical program aims to produce drugs, vaccines and medical devices as countermeasures for CBR threat agents. Specific areas of medical investigation include: prophylaxis, pretreatment, antidotes and therapeutics, personnel and patient decontamination, and medical management of casualties. In the physical sciences area, the focus is on demonstrations of CB defense technologies, including biological detection, chemical detection, and decontamination. The work in this PE is consistent with the Joint Service CB Defense Research, Development, and Acquisition (RDA) Plan. This PE also provides for the conduct of advanced technology development in the areas of real-time sensing, accelerated biological warfare operational awareness, and the restoration of operations following a biological warfare or chemical warfare attack. This program is dedicated to conducting proof-of-principle field demonstrations, test of system-specific technologies to meet specific military needs. Work conducted under this PE transitions to and provides risk reduction for System Integration/ Demonstration (PE 0603884BP/PE 0604384BP) activities.

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Exhibit R-2, RDT&E Budget Item Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY	R-1 ITEM NOMENCLATURE
0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i>	PE 0603384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ATD)</i>
BA 3: <i>Advanced Technology Development (ATD)</i>	

B. Program Change Summary (\$ in Millions)

	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011 Base</u>	<u>FY 2011 OCO</u>	<u>FY 2011 Total</u>
Previous President's Budget	324.769	282.235	0.000	0.000	0.000
Current President's Budget	307.351	299.680	177.113	0.000	177.113
Total Adjustments	-17.418	17.445	177.113	0.000	177.113
• Congressional General Reductions		-1.255			
• Congressional Directed Reductions		0.000			
• Congressional Rescissions	0.000	0.000			
• Congressional Adds		18.700			
• Congressional Directed Transfers		0.000			
• Reprogrammings	-9.223	0.000			
• SBIR/STTR Transfer	-3.650	0.000			
• Other Adjustments	-4.545	0.000	177.113	0.000	177.113

Congressional Add Details (\$ in Millions, and Includes General Reductions)

Project: CI3: *CONGRESSIONAL INTEREST ITEMS (ATD)*

Congressional Add: *Carbon Nanotube Chemical Detector -*

Congressional Add: *Surface Enhanced Infrared Detection of Threats -*

Congressional Add: *Total Perimeter Surveillance (TPS) -*

Congressional Add: *Photo Catalytic Oxidation (PCO) Demonstration for Water Reuse -*

Congressional Add: *Mobile Rapid Response Prototype -*

Congressional Add: *NIDS Automated Bio Agent Identifier -*

Congressional Add: *Portable Rapid Bacterial Warfare Detection Unit -*

Congressional Add: *UCLA High Speed and High Volume Laboratory Network for Infectious Diseases -*

Congressional Add: *Antioxidant Micronutrient Therapeutic Countermeasures for Chemical Agents -*

Congressional Add: *Plant Vaccine Development -*

Congressional Add: *Multi-Purpose Biodefense Immunoarray -*

	<u>FY 2009</u>	<u>FY 2010</u>
	0.791	0.000
	0.987	0.000
	0.989	1.593
	1.373	0.000
	1.082	2.390
	1.000	2.390
	3.156	0.000
	4.862	0.000
	0.792	0.000
	1.582	1.593
	0.792	0.000

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Exhibit R-2, RDT&E Budget Item Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 3: <i>Advanced Technology Development (ATD)</i>	R-1 ITEM NOMENCLATURE PE 0603384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ATD)</i>
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<u>Congressional Add Details (\$ in Millions, and Includes General Reductions)</u>	FY 2009	FY 2010
Congressional Add: <i>Improved CBR Filters -</i>	1.582	0.000
Congressional Add: <i>Acinetobacter Baumannii Research -</i>	1.978	0.000
Congressional Add: <i>Bio Agent Early Warning Detector -</i>	1.978	0.000
Congressional Add: <i>Biological Agent Identifiers -</i>	1.582	0.000
Congressional Add: <i>Eye-Safe Long Range Stand-off System for Detection of Chemical and Biological Weapons -</i>	1.483	0.000
Congressional Add: <i>Mobile Continuous Air Monitor (MCAM) -</i>	1.582	0.000
Congressional Add: <i>Rapid Response Institute -</i>	3.164	0.000
Congressional Add: <i>Liquid Crystal Sensor Technology Research and Development for Force Protection -</i>	2.373	0.000
Congressional Add: <i>Biodefense Vaccine Development and Engineering of Antiviral Peptides -</i>	1.583	0.000
Congressional Add: <i>Center for Advanced Emergency Response -</i>	4.350	0.000
Congressional Add: <i>ViriChip Rapid Virus Detection Systems -</i>	1.582	0.000
Congressional Add: <i>Protective Self-Decontaminating Surfaces -</i>	1.582	0.000
Congressional Add: <i>Contaminated Human Remains Pouch -</i>	1.582	0.000
Congressional Add: <i>Recombinant BChE Formulation Program -</i>	1.582	0.000
Congressional Add: <i>Joint Material Decon System -</i>	1.582	0.000
Congressional Add: <i>Multi-Target Shipping Container Interrogation System Mobile Continuous Air Monitor</i>	0.000	1.593
Congressional Add: <i>Hand-Held Apparatus for Mobile Mapping and Expedited Reporting</i>	0.000	2.788
Congressional Add: <i>Regenerative Chemical Biological Filtration Systems</i>	0.000	2.689
Congressional Add: <i>Unified Management Infrastructure System</i>	0.000	0.797
Congressional Add: <i>Water Purification System for Natural Disasters</i>	0.000	0.797
Congressional Add: <i>CBDP Advanced Development</i>	0.000	1.992
Congressional Add Subtotals for Project: CI3	46.971	18.622

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Exhibit R-2, RDT&E Budget Item Justification: PB 2011 Chemical and Biological Defense Program	DATE: February 2010
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APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 3: <i>Advanced Technology Development (ATD)</i>	R-1 ITEM NOMENCLATURE PE 0603384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ATD)</i>
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Congressional Add Details (\$ in Millions, and Includes General Reductions)

	FY 2009	FY 2010
Congressional Add Totals for all Projects	46.971	18.622

Change Summary Explanation

Funding: N/A - Adjustments less than 10% of total program.

Schedule: N/A

Technical: N/A

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program								DATE: February 2010			
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 3: <i>Advanced Technology Development (ATD)</i>				R-1 ITEM NOMENCLATURE PE 0603384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ATD)</i>				PROJECT CB3: <i>CHEMICAL BIOLOGICAL DEFENSE (ATD)</i>			
COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
CB3: <i>CHEMICAL BIOLOGICAL DEFENSE (ATD)</i>	19.567	25.297	15.410	0.000	15.410	21.450	26.120	36.775	37.148	Continuing	Continuing

A. Mission Description and Budget Item Justification

This project (CB3) demonstrates technology advancements for joint service application in the areas of detection, information systems technology, protection/hazard mitigation, and technology transition efforts. These activities will speed maturing of advanced technologies to reduce risk in system-oriented integration/demonstration efforts. This project also includes efforts dedicated to developing capabilities to protect against Non-Traditional Agents (NTAs). Starting in FY11, all NTA-dedicated research will be re-aligned into specific capability areas within this project in order to ensure a focused effort on this high priority area. Detection focuses on advanced development of technologies from applied research for standoff and point detection and identification of chemical and biological agents. Information systems advanced technology focuses on areas of advanced warning and reporting, hazard prediction and assessment, simulation analysis and planning, and systems performance modeling. Protection and Hazard Mitigation focuses on advanced development of technologies that protect and reduce the chemical/biological threat or hazard to the Warfighter, weapons platforms, and structures. This project also funds advanced development of chemical and biological defense science and technology initiatives and transitions them to advanced development programs in Budget Activities 4 and 5, through prototypes that are evaluated in Advanced Technology Demonstration (ATDs) and Joint Warfighter Experimentation (JWE).

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.	0.000	0.345	0.000	0.000	0.000
2) Decontamination Alternative Processes: Demonstration of non-traditional decontamination technologies and approaches which gain significantly improved effectiveness by complementary application.	1.838	0.000	0.000	0.000	0.000

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 3: <i>Advanced Technology Development (ATD)</i>	R-1 ITEM NOMENCLATURE PE 0603384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ATD)</i>	PROJECT CB3: <i>CHEMICAL BIOLOGICAL DEFENSE (ATD)</i>				
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2009 Accomplishments:</i> Continued efforts to investigate reactive materials and nanotechnology for decontamination processes.</p> <p><i>FY 2010 Plans:</i> Efforts re-aligned to Protection and Hazard Mitigation.</p>						
<p>3) Protection</p> <p>Respiratory/Ocular Protection: Demonstration of design alternatives for chemical and biological air-purifying respirators to provide enhanced protection with lower physiological burden and improved interface with mission equipment.</p> <p><i>FY 2009 Accomplishments:</i> Continued integration of the protective mask designs with developmental helmet systems to provide seamless compatibility of CB protection with ballistic protection, and integration of communication and optical systems. Continued to develop initial high fidelity prototypes for early assessment of human and operational compatibility during the Uniform Integrated Protective Ensemble (UIPE) Demonstration.</p> <p><i>FY 2010 Plans:</i> Efforts re-aligned to Protection and Hazard Mitigation.</p>		1.263	0.000	0.000	0.000	0.000
<p>4) Protection</p> <p>Integrated Ensemble Development: Demonstration of lightweight chemical and biological protective textiles that can be used as an integrated combat duty uniform.</p> <p><i>FY 2009 Accomplishments:</i> Continued integration of the protective mask designs with developmental helmet systems to provide seamless compatibility of chemical and biological protection with ballistic protection, and integration</p>		1.481	0.000	0.000	0.000	0.000

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 3: <i>Advanced Technology Development (ATD)</i>	R-1 ITEM NOMENCLATURE PE 0603384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ATD)</i>	PROJECT CB3: <i>CHEMICAL BIOLOGICAL DEFENSE (ATD)</i>				
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
of communication and optical systems. Continued to develop initial high fidelity prototypes for early assessment of human and operational compatibility during the Uniform Integrated Protective Ensemble (UIPE) Demonstration. <i>FY 2010 Plans:</i> Efforts re-aligned to Protection and Hazard Mitigation.						
5) Protection & Hazard Mitigation Lightweight Integrated Fabric: Demonstration of lightweight chemical and biological protective textiles that can be used as an integrated combat duty uniform. <i>FY 2010 Plans:</i> Develop systems integration of a complete chemical and biological (CB) ensemble that incorporates emerging designs and prototype concepts. Refine concepts for an integrated ensemble that will transition to advanced development programs such as the Uniform Integrated Protective Ensemble (UIPE) and the Individual Protection Advanced Technology Demonstration (IP Demo - see Project TT3, Experimental & Technology Demonstration and Project TT4). Continue limited field trials in a relevant environment. <i>FY 2011 Base Plans:</i> Incorporate lessons from IP Demo and develop final data packages for transition to UIPE and/or Joint Service Lightweight Integrated Suit Technology (JSLIST) programs.		0.000	0.628	0.753	0.000	0.753
6) Protection & Hazard Mitigation Low-Resistance, Low-Profile Filtration: Demonstration of novel filtration media into a lightweight, low-profile, and low-burden individual protective filter, which has enhanced performance against a broader range of challenges that includes toxic industrial chemicals.		0.000	0.635	0.878	0.000	0.878

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p>Logistically Sustainable Air Purification for Collective Protection: Demonstration of chemical and biological air-purification alternative technologies that minimize or eliminate the need for expendable media within acceptable size, weight and power constraints.</p> <p><i>FY 2010 Plans:</i> Initiate breadboard prototypes development of down-selected media-less technologies.</p>						
<p>9) Protection & Hazard Mitigation</p> <p>General Purpose Formulations for Decontamination: Demonstration of improved chemical and biological decontamination formulation that is compatible with the current family of decontamination systems.</p> <p><i>FY 2010 Plans:</i> Complete coupon tests, material compatibility and small item effectiveness evaluations for solid oxidants and green solvent/surfactant systems. Transition to Decontamination Family of Systems program (see BA5, Project DE5).</p>		0.000	0.704	0.000	0.000	0.000
<p>10) Protection & Hazard Mitigation</p> <p>Decontamination System-of-Systems: Demonstration of non-traditional decontamination technologies and approaches which gain significantly improved effectiveness by complementary application.</p> <p><i>FY 2010 Plans:</i> Complete data package for self-decontaminating surfaces. Transition to the Hazard Mitigation for Materials and Equipment Restoration (HaMMER) Advanced Technology Demonstration (see Project TT3, E&TD).</p>		0.000	0.196	0.377	0.000	0.377

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 3: <i>Advanced Technology Development (ATD)</i>	R-1 ITEM NOMENCLATURE PE 0603384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ATD)</i>	PROJECT CB3: <i>CHEMICAL BIOLOGICAL DEFENSE (ATD)</i>				
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
13) Information Systems Technology Battle Space Management: Develop collaborative information management technologies for insertion into the Joint Warning and Reporting Network (JWARN) and Joint Operational Effects Federation (JOEF) acquisition programs. <i>FY 2009 Accomplishments:</i> Transitioned to JWARN the capability to exchange and multi-level fusion of actionable information with real world Command and Control (C2) systems in Department of Defense, Coalition and Homeland Security/Homeland Defense (HLS/HLD) domains. <i>FY 2010 Plans:</i> Battle Space Management efforts re-aligned to Advanced Warning and Reporting.		0.549	0.000	0.000	0.000	0.000
14) Information Systems Technology Advanced Warning and Reporting: Emphasis on developing science and technologies for collaborative information management, fusion of disparate information from multiple sources, environmental databases and modeling, fusion of syndromic/diseases surveillance data, and synthetic environments for model performance evaluation and acquisition decisions. <i>FY 2010 Plans:</i> Transition enhanced version of first-generation building interior Source Term Estimation (STE) and Hazard Refinement (HR) software to the Joint Effects Model (JEM). <i>FY 2011 Base Plans:</i> Transition next-generation outdoor STE, HR, and Sensor Placement Tool (SPT) to advanced development programs (JEM - see BA5 Project IS5). Transition first-generation false alarm reduction capability and first generation rapid STE algorithms to advanced development program (JWARN).		0.000	0.114	1.054	0.000	1.054
15) Information Systems Technology		1.042	1.848	1.961	0.000	1.961

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p>Chemical Biological Defense Program Decision Capability: Develop tools for decision making for consequence management, human knowledge management, and health/human effects modeling including casualty estimation.</p> <p><i>FY 2009 Accomplishments:</i> Verified and incorporated models for casualty estimates for infectious/contagious diseases into JEM. Validated models for predicting effects due to infectious/contagious diseases for Joint Effects Model (JEM) with real-world and simulation data. Completed transition of NATO's AMedP-8 chemical and biological models from NBC CREST to JOEF.</p> <p><i>FY 2010 Plans:</i> CBDP Decision Capability efforts re-aligned to Simulation Analysis and Planning.</p>						
<p>17) Information Systems Technology</p> <p>Chemical and Biological Warfare Effects on Operations: Develop the science behind the modeling and simulation of operations at the strategic, operational and tactical level in a CBRN environment for mobile forces, tactical aircraft, naval operations and fixed sites.</p> <p><i>FY 2009 Accomplishments:</i> Delivered chemical, biological, radiological, and nuclear (CBRN) operational effects methodologies for tactical and theater levels to JOEF. Delivered building interior modeling for JOEF. Completed transition of Agent Fate model to the Joint Effects Model (JEM). Transitioned mobile forces and shipboard models for CB effects on military operations to JOEF. Began validation of decision support tools for CBRN for eventual transition to JOEF.</p> <p><i>FY 2010 Plans:</i> Chemical and Biological Warfare Effects on Operations re-aligned to Simulation Analysis and Planning.</p>		0.821	0.000	0.000	0.000	0.000

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2010 Plans:</i> Complete prototyping a data collection and exchange capability. Complete developing processes and policies for collection and insertion of data into CBRN data backbone efforts.						
20) Detection Detection Capabilities for Non-Traditional Agents: Develop detection technologies for Non-Traditional Agents. <i>FY 2009 Accomplishments:</i> Assessed and demonstrated antibodies assays in handheld format for small chemical molecules. <i>FY 2010 Plans:</i> Continue to develop supporting technologies and protocols to meet the Initial Operating Capabilities of the Next Generation Test Facility at the Edgewood Chemical and Biological Center. <i>FY 2011 Base Plans:</i> All NTA-related efforts re-aligned to the Detection NTA capability area located in this Budget Activity.		2.000	1.964	0.000	0.000	0.000
21) Detection Chemical and Biological Stand-off Technology: Focuses on the detection and identification of chemical and biological threats in near real time at a distance from the detector. Future programs focus on the improvement of algorithms, excitation sources, and detector elements to increase range, reduce false positives, increase sensitivity, and reduce cost. <i>FY 2009 Accomplishments:</i> Completed the fabrication and demonstration technology to meet Joint Biological Standoff Detection System (JBSDS) Increment 2 technology based upon the new information in the infrared		6.611	11.673	0.496	0.000	0.496

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 3: <i>Advanced Technology Development (ATD)</i>	R-1 ITEM NOMENCLATURE PE 0603384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ATD)</i>	PROJECT CB3: <i>CHEMICAL BIOLOGICAL DEFENSE (ATD)</i>
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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p>Technology Transition - Conduct competitive assessments of promising mature technology from outside the Chemical and Biological Defense Program (CBDP) and assist in transition of promising technology efforts.</p> <p><i>FY 2009 Accomplishments:</i> Initiated and completed transition of a miniature, lightweight chemical and biological sensor to JPM-BioDetection from DHS. Initiated transition of the Integrated CB Agent Hazard Mitigation program from the Defense Advanced Research Projects Agency (DARPA) to the United States Army Corps of Engineers through component testing in a laboratory environment. Continued competitive assessment of all mature technology from outside of the CBDP for rapid technology insertion into the capability areas.</p> <p><i>FY 2010 Plans:</i> Continue transition of the Integrated CB Agent Hazard Mitigation with systems and neutralization efficiency testing in a laboratory environment. Continue competitive assessment of all mature technology from outside of the CBDP for rapid technology insertion into the capability areas.</p> <p><i>FY 2011 Base Plans:</i> Complete transition of the Integrated CB Agent Hazard Mitigation with systems and neutralization efficiency testing in an operational environment. Complete assessment and down-select to two or three best technologies that provides the highest enhancements to capabilities.</p>					
Accomplishments/Planned Programs Subtotals	19.567	25.297	15.410	0.000	15.410

C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011 Base</u>	<u>FY 2011 OCO</u>	<u>FY 2011 Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To Complete</u>	<u>Total Cost</u>
• CA4: <i>CONTAMINATION AVOIDANCE (ACD&P)</i>	7.703	40.186	63.347		63.347	9.093	10.754	4.742	3.978	Continuing	Continuing

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C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011</u> <u>Base</u>	<u>FY 2011</u> <u>OCO</u>	<u>FY 2011</u> <u>Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To</u> <u>Complete</u>	<u>Total Cost</u>
• CB2: <i>CHEMICAL BIOLOGICAL DEFENSE (APPLIED RESEARCH)</i>	102.599	110.955	88.897		88.897	100.243	97.979	90.686	91.554	Continuing	Continuing
• DE4: <i>DECONTAMINATION SYSTEMS (ACD&P)</i>	4.822	1.792	7.051		7.051	5.748	1.386	0.000	0.000	Continuing	Continuing
• IS4: <i>INFORMATION SYSTEMS (ACD&P)</i>	0.000	0.000	11.221		11.221	3.404	4.565	4.676	4.741	Continuing	Continuing
• TE3: <i>TEST & EVALUATION (ATD)</i>	25.761	13.307	11.875		11.875	11.267	11.160	0.000	0.000	Continuing	Continuing
• TE4: <i>TEST & EVALUATION (ACD&P)</i>	6.261	28.773	19.304		19.304	11.851	28.035	20.266	21.139	Continuing	Continuing
• TT4: <i>TECHBASE TECHNOLOGY TRANSITION (ACD&P)</i>	17.065	26.649	26.466		26.466	18.564	18.838	19.294	19.563	Continuing	Continuing

D. Acquisition Strategy

N/A

E. Performance Metrics

N/A

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APPROPRIATION/BUDGET ACTIVITY				R-1 ITEM NOMENCLATURE				PROJECT			
0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 3: <i>Advanced Technology Development (ATD)</i>				PE 0603384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ATD)</i>				CI3: <i>CONGRESSIONAL INTEREST ITEMS (ATD)</i>			
COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
CI3: <i>CONGRESSIONAL INTEREST ITEMS (ATD)</i>	46.971	18.622	0.000	0.000	0.000	0.000	0.000	0.000	0.000	Continuing	Continuing

A. Mission Description and Budget Item Justification

The efforts listed in this project include congressional interest programs for FY09 and FY10.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010
Congressional Add: Carbon Nanotube Chemical Detector - <i>FY 2009 Accomplishments:</i> Addressed improvements in sensitivity and selectivity through chemometric/principal component analyses and the development of artificial neural network (ANN) real-time optimum signature selection.	0.791	0.000
Congressional Add: Surface Enhanced Infrared Detection of Threats - <i>FY 2009 Accomplishments:</i> Continued to develop a handheld biological and chemical agent detection device based on surface enhanced infrared detection methods.	0.987	0.000
Congressional Add: Total Perimeter Surveillance (TPS) - <i>FY 2009 Accomplishments:</i> Demonstrated a prototype of the system.	0.989	1.593

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B. Accomplishments/Planned Program (\$ in Millions)		
	FY 2009	FY 2010
<i>FY 2010 Plans:</i> Continuation of FY09 research.		
Congressional Add: Photo Catalytic Oxidation (PCO) Demonstration for Water Reuse - <i>FY 2009 Accomplishments:</i> Continued research to determine the water purification unit's performance in the removal of high threat CBRN agents and TICs.	1.373	0.000
Congressional Add: Mobile Rapid Response Prototype - <i>FY 2009 Accomplishments:</i> Continued the partnership of Hackensack University Medical Center with the Defense Threat Reduction Agency (DTRA), the Chemical Biological & Radiological Technology Alliance. <i>FY 2010 Plans:</i> Continuation of FY09 research.	1.082	2.390
Congressional Add: NIDS Automated Bio Agent Identifier - <i>FY 2009 Accomplishments:</i> Continued research begun in FY08. <i>FY 2010 Plans:</i> Conduct research for the development of multiplex handheld immunoassay tickets that are both human visually and machine read.	1.000	2.390
Congressional Add: Portable Rapid Bacterial Warfare Detection Unit -	3.156	0.000

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B. Accomplishments/Planned Program (\$ in Millions)		
	FY 2009	FY 2010
<i>FY 2009 Accomplishments:</i> Developed a field deployable system based on IR spectroscopy.		
Congressional Add: UCLA High Speed and High Volume Laboratory Network for Infectious Diseases - <i>FY 2009 Accomplishments:</i> Expanded capability to include other biothreat agents, including bacterial and/or viruses (dual-use).	4.862	0.000
Congressional Add: Antioxidant Micronutrient Therapeutic Countermeasures for Chemical Agents - <i>FY 2009 Accomplishments:</i> Tested the hypothesis that a mixture of antioxidants before and after exposure to sulfur mustard may increase percent survival and survival time by decreasing oxidative damage and inflammation.	0.792	0.000
Congressional Add: Plant Vaccine Development - <i>FY 2009 Accomplishments:</i> Produced vaccine lots under cGMP and evaluated safety and toxicity and confirmed protective efficacy of identified dual agent vaccines. Developed technology transfer and implementation programs. <i>FY 2010 Plans:</i> Continuation of FY09 Research.	1.582	1.593
Congressional Add: Multi-Purpose Biodefense Immunoarray - <i>FY 2009 Accomplishments:</i> Continued research from FY08.	0.792	0.000

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B. Accomplishments/Planned Program (\$ in Millions)			
		FY 2009	FY 2010
Congressional Add: Improved CBR Filters - <i>FY 2009 Accomplishments:</i> Initiated engineering phase with the goal of developing final design configurations that can be easily incorporated into new and existing filtration systems.		1.582	0.000
Congressional Add: Acinetobacter Baumannii Research - <i>FY 2009 Accomplishments:</i> Continued the preclinical development of these agents by developing improved syntheses techniques.		1.978	0.000
Congressional Add: Bio Agent Early Warning Detector - <i>FY 2009 Accomplishments:</i> Conducted advanced development of a standoff bio agent detection system.		1.978	0.000
Congressional Add: Biological Agent Identifiers - <i>FY 2009 Accomplishments:</i> Continued industry research into biological agent identifiers without wet reagents.		1.582	0.000
Congressional Add: Eye-Safe Long Range Stand-off System for Detection of Chemical and Biological Weapons - <i>FY 2009 Accomplishments:</i> Continued research for eye-safe, laser based standoff Chem/Bio detection systems.		1.483	0.000
		1.582	0.000

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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010
Congressional Add: Mobile Continuous Air Monitor (MCAM) - <i>FY 2009 Accomplishments:</i> Continued research for a portable continuous monitor for biodetection.		
Congressional Add: Rapid Response Institute - <i>FY 2009 Accomplishments:</i> Addressed technology related to quickly responding to chemical or biological situation.	3.164	0.000
Congressional Add: Liquid Crystal Sensor Technology Research and Development for Force Protection - <i>FY 2009 Accomplishments:</i> Continued development of a passively operated sensor that rapidly detects toxins in the immediate environment.	2.373	0.000
Congressional Add: Biodefense Vaccine Development and Engineering of Antiviral Peptides - <i>FY 2009 Accomplishments:</i> Performed vaccine development pertaining to antiviral countermeasures.	1.583	0.000
Congressional Add: Center for Advanced Emergency Response - <i>FY 2009 Accomplishments:</i> Continued development of emergency medical response training program for consequence management of chemical or biological events.	4.350	0.000
Congressional Add: ViriChip Rapid Virus Detection Systems -	1.582	0.000

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B. Accomplishments/Planned Program (\$ in Millions)		
	FY 2009	FY 2010
<i>FY 2009 Accomplishments:</i> Researched on the use of nanoscience technology for a virus detection system.		
Congressional Add: Protective Self-Decontaminating Surfaces - <i>FY 2009 Accomplishments:</i> Provided immediate on-site protection with multi-threat applicability to instantly neutralize chemical agents and kill a number of microbial entities.	1.582	0.000
Congressional Add: Contaminated Human Remains Pouch - <i>FY 2009 Accomplishments:</i> Conducted prototype development activities to test a contaminated human remains transportable container.	1.582	0.000
Congressional Add: Recombinant BChE Formulation Program - <i>FY 2009 Accomplishments:</i> Conducted medical countermeasure technology development.	1.582	0.000
Congressional Add: Joint Material Decon System - <i>FY 2009 Accomplishments:</i> Addressed Reactive Overlay and Removable CBRN Coatings.	1.582	0.000
Congressional Add: Multi-Target Shipping Container Interrogation System Mobile Continuous Air Monitor	0.000	1.593

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B. Accomplishments/Planned Program (\$ in Millions)		
	FY 2009	FY 2010
<i>FY 2010 Plans:</i> Develop air monitoring system for shipping containers.		
Congressional Add: Hand-Held Apparatus for Mobile Mapping and Expedited Reporting <i>FY 2010 Plans:</i> Develop a tool that enables a rapid, accurate, efficient, low-cost, collection, analysis and dissemination of digital data from multiple sensor suites and rapid reporting for improved situational awareness.	0.000	2.788
Congressional Add: Regenerative Chemical Biological Filtration Systems <i>FY 2010 Plans:</i> Continuation of research funded in FY08 for a regenerative filtration system to reduce costs and provide protection against all chemical warfare agents for military personnel, critical equipment, and strategic facilities.	0.000	2.689
Congressional Add: Unified Management Infrastructure System <i>FY 2010 Plans:</i> Develop a secure communication platform to meet military needs in a chemical biological environment, protecting soldiers and first responders on the battlefield using secure mobile communication systems by simultaneously providing what is currently unprecedented: real-time, accurate monitoring of the military's communication devices.	0.000	0.797
Congressional Add: Water Purification System for Natural Disasters <i>FY 2010 Plans:</i> Develop a water purification system.	0.000	0.797

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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010
Congressional Add: CBDP Advanced Development <i>FY 2010 Plans:</i> Advanced Development	0.000	1.992
Congressional Adds Subtotals	46.971	18.622

C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011</u> <u>Base</u>	<u>FY 2011</u> <u>OCO</u>	<u>FY 2011</u> <u>Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To</u> <u>Complete</u>	<u>Total Cost</u>
• C11: <i>CONGRESSIONAL INTEREST ITEMS (BASIC RESEARCH)</i>	8.090	20.036	0.000		0.000	0.000	0.000	0.000	0.000	Continuing	Continuing
• C12: <i>CONGRESSIONAL INTEREST ITEMS (APPLIED RESEARCH)</i>	42.714	16.630	0.000		0.000	0.000	0.000	0.000	0.000	Continuing	Continuing

D. Acquisition Strategy

N/A

E. Performance Metrics

N/A

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0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 3: <i>Advanced Technology Development (ATD)</i>				PE 0603384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ATD)</i>				TB3: <i>MEDICAL BIOLOGICAL DEFENSE (ATD)</i>			
COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
TB3: <i>MEDICAL BIOLOGICAL DEFENSE (ATD)</i>	180.425	203.723	115.233	0.000	115.233	125.666	109.737	115.049	117.289	Continuing	Continuing

A. Mission Description and Budget Item Justification

This project (TB3) funds preclinical development of vaccines, therapeutic drugs, and diagnostic capabilities to provide safe and effective medical defense against validated biological threat agents including bacteria, toxins, and viruses. Innovative biotechnology approaches to advance medical systems designed to rapidly identify, diagnose, prevent, and treat disease due to exposure to biological threat agents will be evaluated. Entry of candidate vaccines, therapeutics, and diagnostic technologies into advanced development is facilitated by the development of technical data packages that support the Food and Drug Administration (FDA) Investigational New Drug (IND) licensure processes, DoD acquisition regulations, and the oversight of Phase 1 clinical trials in accordance with FDA guidelines. Categories of this project include biological defense capability areas such as Pretreatments, Diagnostics, and Therapeutics. Pretreatment efforts conduct research and development (R&D) of promising vaccines, medications, and technologies provided prior to potential exposure to biological agents. The goal is to reduce or to entirely prevent adverse effects of exposure. Diagnostic efforts are aimed at screening procedures and analytical methods to verify exposure and determine the effects of exposure to biological warfare (BW) agents. Therapeutic efforts provide medical solutions to sustain and protect the Warfighter in biological environments. Specifically, therapeutic efforts are aimed at developing medical countermeasures to treat exposure to biological threats such as bacterial (plague, anthrax, glanders), viral (smallpox, encephalitic alphaviruses), and toxin (ricin, botulinum neurotoxin, staphylococcal enterotoxin) agents.

This project also includes efforts such as the Transformational Medical Technologies Initiative (TMTI). The Transformational Medical Technologies Initiative (TMTI) was launched to respond to the threat of emerging or intentionally bioengineered biological threats. TMTI's mission is to protect the Warfighter from genetically engineered biological threats by providing a rapid response capability from identification of pathogens to the delivery of medical countermeasures. This mission is accomplished through two main efforts: 1) developing broad spectrum (multi-agent) therapeutics against BW agents (e.g, one drug that treats multiple agents); and 2) developing platform technologies to assist in the rapid development of medical countermeasures (MCMs) in response to BW agents (e.g, developing new and innovative ways to mass produce drugs in the event of a biological incident).

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) SBIR	0.000	2.637	0.000	0.000	0.000

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2010 Plans:</i> Small Business Innovative Research.						
2) Diagnostics Diagnostic Technologies: Development and verification of rapid, sensitive and specific tests for the identification of Biological Warfare Agents (BWAs) and their expressed toxins in biological fluids of Warfighters for the diagnosis of exposure/infection. Discovery of biomarkers of response to exposure. Evaluation of next generation diagnostic technologies including portable instrument platforms, highly parallel and informative testing formats, and nanotechnology applications. <i>FY 2009 Accomplishments:</i> Transitioned two candidates for a next generation diagnostic device to the advanced developer. Continued to utilize the decision matrix to identify and evaluate new technologies more effective for diagnosing exposure to bio-threat agents. Validated real time PCR assays identifying genes responsible for antibiotic resistance in bio-threat agents. Performed advanced assessment on the use of biosynthetic (recombinant) reagents on existing systems and improved test assays utilizing new technologies and approaches that enhance diagnosis of early exposure to BWAs. <i>FY 2010 Plans:</i> Continue development of two additional candidates for a next generation diagnostic device. Develop an automated, prototype polymerase chain reaction system on microarray cartridge using light emitting chemical-based (or other sensitive signal-amplified) technology. Continue to refine and transition strain test panels for viral specificity (inclusivity and exclusivity) characterization. Characterize assay specificity to ensure assays consistently identify the intended target but not related targets. Use highly parallel and informative microarray screening techniques with thoroughly characterized affinity reagents for the discovery of novel biomarkers of host response as targets for assay development. Develop and verify assays as per standardized processes. Transition pilot production protocols for biosynthetic (recombinant) antigen production for bacterial BWAs. Maintain an animal tissue bank		9.021	11.307	9.845	0.000	9.845

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p>for validation of assay performance and as correlate reference materials from animal BWA exposure studies. Develop and verify single domain biosynthetic (recombinant) antibodies to bacterial and viral BWA targets. Investigate methods of stabilization of BWA biomarkers in clinical samples to extend transport and limit cold chain requirements.</p> <p><i>FY 2011 Base Plans:</i> Use decision-based matrix and technology evaluation centers to transition two Technology Readiness Reviews on candidate diagnostic platforms to advanced development programs. Develop atlas/database of phenotypic and genotypic characteristics of relevant BWA bacterial strains. Demonstrate the utility of high informatic content screen-characterized affinity reagents in the discovery of novel biomarkers as targets for assay development. Develop standard methods/protocols for rapid sequencing directly from clinical matrices. Apply bioinformatic and computational methods to verify the utility of host response signatures for pre-symptomatic diagnostic assays. Transition candidate transport media/preservatives and protocols for clinical sample processing. Evaluate developed global-virus and global-microbial microarrays for promising multiplexing and identification of BWAs. Develop and verify production scale-up protocols for single domain biosynthetic (recombinant) antibodies to bacterial and viral BWA targets.</p>						
<p>3) Pretreatments</p> <p>Vaccine Research Support: Assess the effectiveness of candidate vaccines in animal models and perform preliminary evaluations of safety and duration of protective immunity.</p> <p><i>FY 2009 Accomplishments:</i> Further characterized safety, toxicity, and immunity duration studies in animals for filovirus vaccines. Optimized dose, route, and regimen for maximum effectiveness. Assessed alphavirus and filovirus vaccines for issues of vaccine interference. Conducted stability and toxicity studies for lead alphavirus vaccine candidates. Completed stability and toxicity studies for toxin vaccines, prepared production lots, and begin Investigational New Drug (IND) application preparation for Food and Drug</p>		8.128	0.000	0.000	0.000	0.000

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
Administration (FDA) evaluation. Analyzed effectiveness, duration of immunity, and dosing regimens of second-generation vaccine against bacterial pathogens (including anthrax, plague, and tularensis). <i>FY 2010 Plans:</i> Vaccine Research Support efforts re-aligned to Bacterial/Toxin and Viral Vaccines.						
4) Pretreatments Multiagent Vaccine Platforms: Evaluates the safety and effectiveness of vaccine platforms for immunization against multiple biothreat agents. <i>FY 2009 Accomplishments:</i> Evaluated safety and effectiveness of multi-agent vaccines (e.g., anthrax/plague/melioidosis); completed studies to determine interference between vaccine components and the immune response; conducted immunity duration studies. Down-selected multiagent vaccine platforms, determine dosage, and route of administration. <i>FY 2010 Plans:</i> Multi-agent Vaccine efforts will be re-aligned to Vaccine Platforms and Research Tools.		3.500	0.000	0.000	0.000	0.000
5) Pretreatments Bacterial/Toxin Vaccines: Evaluates the best single agent bacterial and toxin vaccines for effectiveness against aerosol challenge in large animal models. <i>FY 2010 Plans:</i> Plan, prepare and conduct Phase I clinical trial with the Ricin vaccine. <i>FY 2011 Base Plans:</i> Complete the Phase I clinical trial with the Ricin Vaccine.		0.000	0.984	0.937	0.000	0.937

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APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 3: <i>Advanced Technology Development (ATD)</i>		R-1 ITEM NOMENCLATURE PE 0603384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ATD)</i>		PROJECT TB3: <i>MEDICAL BIOLOGICAL DEFENSE (ATD)</i>		
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
6) Pretreatments		0.000	16.372	10.304	0.000	10.304
<p>Viral Vaccines: Evaluates the best vaccine candidates for Alphaviruses and Filoviruses for effectiveness and duration of protective immune response against aerosol challenge in large animal models. Animal models will be developed to support FDA licensure of mature vaccine candidates.</p> <p><i>FY 2010 Plans:</i> Initiate studies to develop/validate animal models for VEE, EEE, and WEE vaccines, as well as for filovirus vaccines, to fulfill future FDA animal rule requirements necessary for vaccine licensure. Test chemically inactivated and deoxyribonucleic acid (DNA) vaccine candidates against VEE, EEE, and WEE for effectiveness against aerosol delivered doses in animals. Conduct dose, schedule, and aerosol challenge studies in animals with Ebola vaccine candidates. Transition two Marburg virus vaccine candidates to advanced development programs, and determine protection duration studies on these two candidates. Conduct studies to further evaluate the effectiveness of combining the individual filoviruses (i.e., Ebola Sudan, Ebola Zaire, Ebola Uganda, and Marburg Angola) vaccines into one multi-agent vaccine. Conduct studies to further evaluate the effectiveness of combining the individual alphavirus (i.e., VEE, EEE, and WEE) vaccines into one multi-agent vaccine.</p> <p><i>FY 2011 Base Plans:</i> Complete duration studies with the vaccine components against Marburg that transitioned to the advanced development program in FY10. Complete aerosol efficacy studies for the Ebola Zaire and Ebola Sudan vaccine components in non-human primates. Transition the Ebola vaccine components to the advanced development program to combine with the Marburg vaccine component. Determine duration of protection elicited by the Ebola vaccine components. Optimize the dose and immunization schedule to ensure effectiveness of the individual components of the filovirus vaccine when co-administered as a mixture. Complete aerosol efficacy studies of DNA-based vaccines and chemically inactivated/attenuated vaccines against the alphaviruses. Optimize dosing regimens to ensure effectiveness when co-administering the alphavirus vaccine components. Continue the development of animals models for alphaviruses (EEE and WEE), and filoviruses (Ebola Sudan, Ebola Zaire,</p>						

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
responses elicited by vaccines and/or pathogens of interest, and compare those results to animal studies. Evaluate the safety and immune stimulating capability of mature Filovirus and Alphavirus vaccine candidates in humans by using the MIMIC technology, to support these candidates moving forward into phase I clinical studies by the advanced development program. Conduct pre-formulation studies to produce a thermo-stable, spray-dried formulation of the virus-like particle based Marburg vaccine candidate.						
8) Therapeutics Therapy for Ebola and Marburg Virus Infections: Identifies, optimizes and evaluates potential therapeutic candidates effective against Filovirus infection including Ebola and Marburg Viruses. <i>FY 2009 Accomplishments:</i> Completed FDA required studies to support the preclinical development and characterization of other leading therapeutic technologies against the Ebola virus and Marburg virus.		5.302	0.000	0.000	0.000	0.000
9) Therapeutics Viral Therapeutics: Identifies, optimizes and evaluates potential therapeutic candidates effective against designated viral threat agents. <i>FY 2009 Accomplishments:</i> Continued studies to support FDA submissions, milestone approval, and product transition to advanced development programs. Performed FDA required non-human primate studies necessary to complete the development of two oral therapeutics for orthopox viral infection. <i>FY 2010 Plans:</i> Conduct non-human primate studies to determine if anti-inflammatory and anti-thrombotic host factors can be used therapeutically to produce a restorative effect on the blood vessel walls and increase survival from filovirus infection. Conduct remaining FDA required non-human primate studies		5.567	9.493	9.519	0.000	9.519

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p>necessary to complete the development of oral therapeutics for orthopox viral infection. Evaluate the efficacy of administering post-exposure therapeutic vaccine in conjunction with therapies that stop blood clotting in animals infected with filovirus. Continue animal studies to support FDA submissions, milestone approval, and product transition to advanced development.</p> <p><i>FY 2011 Base Plans:</i> Conduct remaining non-human primate studies required for licensure of ST-246, a low-molecular-weight compound that is active against multiple orthopoxviruses. Conduct toxicology studies and analyze efficacy of optimized lead compounds against alphavirus infection in murine and non-human primate challenge models. Characterize the clinical manifestations and virologic/immunologic parameters of human monkeypox. Determine the effectiveness of pan-alphavirus capsid assembly inhibitors in animal models.</p>						
<p>10) Therapeutics</p> <p>Bacterial Therapeutics: Identifies, optimizes, and evaluates potential therapeutic compounds effective against bacterial threat agents.</p> <p><i>FY 2009 Accomplishments:</i> Tested and evaluated FDA approved antibiotics for efficacy against aerosol exposure to bacterial threat agents in non-human primate models of plague. Initiated advanced safety and effectiveness studies for a new single domain antibody that is smaller than conventional antibodies against plague.</p> <p><i>FY 2010 Plans:</i> Test and evaluate the effectiveness of commercially available antibiotics against animals exposed to aerosol versions of plague and tularemia. Determine antibiotic susceptibility profiles for Yersinia pestis and Francisella tularensis in the laboratory.</p>		2.447	2.656	2.700	0.000	2.700

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2011 Base Plans:</i> Determine the effectiveness of commercially available antibiotics against Francisella tularensis in relevant animal infection models.						
11) Therapeutics Toxin Therapeutics: Identifies, optimizes and evaluates potential therapeutic candidates effective against biological toxin threat agents. <i>FY 2009 Accomplishments:</i> Continued optimization and structural activity relationship studies for BoNT small molecule therapeutics to achieve improved pharmacological properties. Tested intraneuronal delivery of small molecules using prototype therapeutic delivery system. Evaluated immune modifying compounds for pre and post-exposure therapy for Staphylococcal Enterotoxin B (SEB) intoxication. <i>FY 2010 Plans:</i> Initiate work to develop antitoxin preparation for Ricin and Staphylococcal Enterotoxin B (SEB). Define the therapeutic parameters for Ricin and SEB therapeutic. Test candidate BoNT small molecule therapeutics in animal challenge models. Perform advanced animal testing on small molecules that are protective against a lethal challenge of SEB in relevant animal models. <i>FY 2011 Base Plans:</i> Test and evaluate FDA approved immunomodulating drugs against exposure to SEB. Develop and determine the therapeutic window of opportunity for novel inhibitors of SEB pathogenesis. Determine initial safety profile and conduct genotoxicity studies for BoNT inhibitors with the goal of improving physiochemical properties and mitigating product liabilities through the use of medicinal chemistry. Conduct pre- and post-challenge of efficacy studies of optimized BoNT inhibitors in mice. Evaluate efficacy of BoNT lead inhibitors using a targeted delivery system in mice.		1.683	1.475	1.500	0.000	1.500

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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p>12) Transformational Medical Technologies Initiative</p> <p>Multiagent (Broad Spectrum) Medical Countermeasures: Focuses on the initiation and completion of multiple preclinical studies for each new drug, to include safety, toxicity, efficacy, and scalability work. The ability to formulate good manufacturing pilot lots and further mature promising drug candidates will be the focus of activities in this capability area. The preclinical drug discovery process culminates in the submission of an Investigational New Drug (IND) application to the Food and Drug Administration (FDA), which conducts reviews and approves new drug candidates. Estimated attrition from preclinical phase to Phase I clinical studies is approximately 50%, thus not all drugs will survive the transition between preclinical development and Phase I studies.</p> <p><i>FY 2009 Accomplishments:</i> Continued to identify potential IND candidate drugs for development. Completed pre-clinical research necessary to submit up to ten additional applications for an IND with the FDA. Accelerated drug discovery efforts, incorporating new technology to expand the number of potential drug compounds suitable for advanced development. Implemented use of the previously validated transgenic and other animal model systems to replicate human disease and disease response pathways. Began implementation of test platforms for drug discovery, development, and manufacturing technologies. Continued investigating the use of existing of FDA-approved drugs to enhance effectiveness of current BW agent countermeasures.</p> <p><i>FY 2010 Plans:</i> Continue to identify potential IND candidate drugs for development. Complete pre-clinical research necessary to submit up to seven additional applications for an IND with the FDA. Upon submission of an IND to the FDA for further evaluation, DoD Milestone A decisions will take place. Downselect contract performers who have had their IND applications accepted by the FDA. Initiate planning for Phase 1 clinical trials and other studies necessary to support advanced development efforts toward an New Drug Application (NDA) with the FDA. Continue investigating use of existing of FDA-approved drugs to enhance effectiveness of current BW agent countermeasures.</p>	144.777	124.132	63.135	0.000	63.135

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p>drug discovery and development platform technologies. Accelerate effort to develop and scale-up new rapid manufacturing platform technologies for biological drugs. Development efforts will bring these technologies into compliance with FDA current good manufacturing practices (cGMP) and quality requirements. Generate Technology Development Strategies that will assist in the development of a roadmap to support efforts that transition to engineering, manufacturing, and development efforts in Budget Activities 4 and 5. Begin integration of stand-alone platforms into capabilities that can be demonstrated as a system. Validate test platforms for drug discovery, development and manufacturing technologies that allow the incorporation of medical countermeasure technologies into the TMTI rapid response capability. Support computer models to advance/enhance drug design. High throughput screening assays and technologies and novel platforms for target identification will also be investigated.</p> <p><i>FY 2011 Base Plans:</i> Continue integration of standalone platforms into capabilities that can be demonstrated as a system. Continue the development of rapid drug discovery and development platform technologies. Integrate the entire system using a robust bioinformatics capability, and validate the integrated bioinformatics platform. Continue to mature and accelerate manufacturing platform technologies for biological drugs to comply with regulatory guidelines. Support compliance and quality measures that are mandatory for future FDA submissions. Continue to integrate pathogen characterization, target identification, countermeasure discovery and countermeasure evaluation platform areas into a rapid response capability supported by a centralized bioinformatics capability that ties together geographically separated performers from government agencies, industry and academia.</p>						
Accomplishments/Planned Programs Subtotals		180.425	203.723	115.233	0.000	115.233

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C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011</u> <u>Base</u>	<u>FY 2011</u> <u>OCO</u>	<u>FY 2011</u> <u>Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To</u> <u>Complete</u>	<u>Total Cost</u>
• MB4: <i>MEDICAL BIOLOGICAL DEFENSE (ACD&P)</i>	7.910	102.437	136.975		136.975	130.718	131.347	115.985	113.566	Continuing	Continuing
• MB5: <i>MEDICAL BIOLOGICAL DEFENSE (SDD)</i>	87.676	57.558	141.680		141.680	161.732	159.144	141.481	111.671	Continuing	Continuing
• TB2: <i>MEDICAL BIOLOGICAL DEFENSE (APPLIED RESEARCH)</i>	50.485	53.930	43.858		43.858	50.866	51.077	51.051	51.959	Continuing	Continuing

D. Acquisition Strategy

N/A

E. Performance Metrics

N/A

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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
TC3: <i>MEDICAL CHEMICAL DEFENSE (ATD)</i>	21.641	28.971	29.134	0.000	29.134	30.401	30.546	31.356	31.877	Continuing	Continuing

A. Mission Description and Budget Item Justification

This project (TC3) supports the advanced development of medical countermeasures to include prophylaxes, pretreatments, antidotes, skin decontaminants and therapeutic drugs against identified and emerging chemical warfare threat agents. Analytical stability studies, safety and efficacy screening, and preclinical toxicology studies are performed prior to full-scale development of promising pretreatment or treatment drug compounds. Entry of candidate pretreatment/prophylaxes, therapeutics, and diagnostic technologies into advanced development (i.e., efforts funded in Budget Activities 4 and 5) is facilitated by the development of technical data packages that support the Food and Drug Administration (FDA) Investigational New Drug (IND) application and licensure processes, as well as Department of Defense (DoD) acquisition regulations. Categories for this project include Pretreatments, Diagnostics, and Therapeutics to address Chemical Warfare Agent (CWA) and Non-Traditional Agents (NTAs) exposure. Starting in FY11, all NTA-dedicated research will be re-aligned into specific capability areas within this project in order to ensure a focused effort on this high priority area.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.	0.000	0.373	0.000	0.000	0.000
2) Diagnostics Diagnostic Technologies: Focuses on state-of-the-art laboratory/fieldable methods that detect exposure to chemical warfare agents (CWA) (e.g., nerve agents and vesicants) in clinical samples. It also targets the identification of biomolecular targets that can be leveraged as analytical methodologies, as well as, laboratory and animal studies characterizing time-course and longevity of a particular analyte/ biomarker.	0.701	1.436	0.226	0.000	0.226

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B. Accomplishments/Planned Program (\$ in Millions)								
				FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2009 Accomplishments:</i> Concluded validation of the optimized sulfur mustard blood protein assay. Initiated validation of the urine byproduct assay. Concluded metabolic profile study and conduct data analysis. Completed validation of procedure to assess the presence of chemical warfare analytes from hair samples.</p> <p><i>FY 2010 Plans:</i> Further development of improved reactivation and solvent-free extraction methodologies for definitive CWA byproduct identification. Determine windows of opportunity for biomarker identification and subsequent therapeutic intervention for CWA in laboratory and animal models.</p> <p><i>FY 2011 Base Plans:</i> Optimize the methodology for solvent free extraction of CWA mixtures. Complete blood and urine assay development for CWA exposure. Complete validation of fluoride regeneration method in plasma/blood/RBCs with solid phase extraction for nerve agents. All NTA-specific efforts re-aligned to the Chemical Diagnostics NTA capability area within this Budget Activity.</p>								
3) Chem Diagnostics NTA <i>FY 2011 Base Plans:</i> Continue evaluation of mature technologies that can quickly diagnose NTA exposure before symptoms appear and determine the type of agent.				0.000	0.000	0.400	0.000	0.400
4) Pretreatments Nerve Agent, Bioscavengers: Develop pretreatments that provide protection against all organophosphorous nerve agents. Bioscavengers should have the ability to rapidly bind and detoxify nerve agents, and have broad binding specificity and high catalytic efficiency for the destruction of agents. One molecule of catalytic bioscavenger should be capable of detoxifying numerous molecules				6.636	7.811	7.861	0.000	7.861

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B. Accomplishments/Planned Program (\$ in Millions)								
				FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2009 Accomplishments:</i> Initiated animal studies to determine long term effects of down-selected wound healing products and blister agents, in coordination with the advanced developer.</p> <p><i>FY 2010 Plans:</i> Evaluate commercial off-the-shelf irrigation systems for treatment of CWA exposure in the laboratory and animals. Continue animal studies to examine long-term effects of wound healing products. Down-select newly identified therapeutics with potential for treating mustard agent-induced ocular injury. Begin efficacy testing in compliance with FDA regulations for ocular administration.</p> <p><i>FY 2011 Base Plans:</i> Continue to evaluate the effectiveness of various cell-based approaches to facilitate blister agent wound healing in skin and eye. Begin advanced studies focused on down-selecting wound healing products found to be most effective for transition. Continue to assess in animals whether bioengineering and molecular biology approaches may be used to treat blister agent skin and eye injury. Initiate the development of an approach to decontaminate CWAs in penetrating wounds.</p>								
6) Therapeutics				6.421	13.240	13.137	0.000	13.137
<p>Neurologic: Focuses on therapeutic strategies to effectively minimize neurologic injuries resulting from exposure to CWAs. This effort involves the development of neuroprotectants, anticonvulsants, and improved neurotransmitter restorers. Supports eventual FDA licensure of new compounds or new indications for licensed products for use in the treatment of chemical warfare casualties.</p> <p><i>FY 2009 Accomplishments:</i> Accelerated efforts to evaluate novel and FDA approved anticonvulsants, neuroprotectants, anti-epileptics, and receptor competitors and neutralizing agents for neuroprotective activity against nerve agents in animal models according to FDA guidelines.</p>								

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B. Accomplishments/Planned Program (\$ in Millions)								
				FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2010 Plans:</i> Test broad-spectrum reactivators in one or more animal models, with a focus on requirements to support FDA submissions under the animal rule. Initiate safety/side effect/dosing and the body's effects on the drug evaluation of new compounds. Continue to evaluate novel and FDA-approved anticonvulsants, neuroprotectants, anti-epileptics, and receptor competitors and neutralizing agents for neuroprotective activity against nerve agents in animal models.</p> <p><i>FY 2011 Base Plans:</i> Continue to evaluate, in animals, novel compounds and FDA-approved drugs not yet evaluated for efficacy against nerve agents. These potential compounds include anticholinergics, neuroprotectants, anticonvulsants, and improved reactivators. Continue efficacy testing on candidates that are designed to support eventual FDA licensure. Continue development of animals models related to nerve exposure with emphasis on FDA animal rule approval.</p>								
7) Therapeutics Medical Toxicology (Non-Traditional Agents (NTAs) and Other agents): Investigates common mechanisms of agent injury. Determines the toxic effects of agents by probable routes of field exposure as well as standard experimental routes. Physiological parameters and pathological assessment will be used to establish the general mode and mechanism(s) of toxicity. <i>FY 2009 Accomplishments:</i> Developed, validated, and completed practical clinical strategies to aid in management of NTA casualties.				2.950	0.000	0.000	0.000	0.000
8) Therapeutics CWA Operational Exposure Hazard Assessment Research: Supports FDA licensure of new compounds or new indications for licensed products for use in the treatment of chemical warfare casualties.				1.000	0.000	0.000	0.000	0.000

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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p>Non Traditional Agents (NTAs): Determines the toxic effects of agents by probable routes of field exposure and refines standard experimental routes. Physiological parameters and pathological assessment will be used to establish the general mode and mechanisms of toxicity.</p> <p><i>FY 2010 Plans:</i> Develop and evaluate novel and FDA licensed products as post-exposure therapeutics against NTA poisoning in advanced animal models.</p> <p><i>FY 2011 Base Plans:</i> Complete characterization of a novel therapeutic for manufacturability and pharmacology. Establish formulation for safety testing and stability.</p>					
Accomplishments/Planned Programs Subtotals	21.641	28.971	29.134	0.000	29.134

C. Other Program Funding Summary (\$ in Millions)

Line Item	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	FY 2012	FY 2013	FY 2014	FY 2015	Cost To Complete	Total Cost
• MC4: <i>MEDICAL CHEMICAL DEFENSE (ACD&P)</i>	19.365	9.438	0.000		0.000	2.973	3.661	5.035	14.670	Continuing	Continuing
• MC5: <i>MEDICAL CHEMICAL DEFENSE (SDD)</i>	14.203	14.027	51.856		51.856	47.835	28.771	12.122	8.171	Continuing	Continuing
• TC2: <i>MEDICAL CHEMICAL DEFENSE (APPLIED RESEARCH)</i>	35.008	40.418	33.648		33.648	36.327	36.500	37.475	38.150	Continuing	Continuing

D. Acquisition Strategy

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E. Performance Metrics

N/A

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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
TE3: <i>TEST & EVALUATION (ATD)</i>	25.761	13.307	11.875	0.000	11.875	11.267	11.160	0.000	0.000	Continuing	Continuing

A. Mission Description and Budget Item Justification

This project (TE3) supports the development of test and evaluation methodologies and protocols as new science and technology efforts are discovered and transitioned to advanced development programs. It includes methodology development for chemical and biological defense test and evaluation capabilities, with an emphasis on Non Traditional Agents (NTAs). These methodologies support development testing and operational testing with regard to advanced development programs that have unique chemical and biological defense requirements. These new methodologies and testing capabilities include the development of protocol and standards for use of chemical and biological simulants. Starting in FY11, all NTA-dedicated research will be re-aligned into specific capability areas within this project.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.	0.000	0.176	0.000	0.000	0.000
2) Test and Evaluation (T&E) NTA Develops test and evaluation technologies and processes in support of NTA activities. <i>FY 2011 Base Plans:</i> Conduct facility design efforts by conducting large particle dissemination development and proof of principle tests with several agents. Complete testing regarding the safety of unprotected personnel using the chamber after decontamination.	0.000	0.000	2.000	0.000	2.000
3) Test and Evaluation (T&E)	6.643	5.896	2.784	0.000	2.784

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B. Accomplishments/Planned Program (\$ in Millions)								
				FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2009 Accomplishments:</i> Completed development of collective protection shelter systems test and evaluation standards, Toxic Industrial Chemicals (TIC), and battlefield contaminant standards for Individual Protection Equipment (IPE) and Collective Protection (COLPRO). Completed standard procedures for IPE Assessment. Completed real-time sampling/detector system swatch test methodology for use in Chemical and Biological Agent Resistance Test System (CBARTS) test methodology standards and guidance for air purification technologies, IPE field operations effects standard, and IPE air flow mapping.</p> <p><i>FY 2010 Plans:</i> Initiate methodology/source data effort to simulate IP durability test in laboratory and relationship to field durability.</p> <p><i>FY 2011 Base Plans:</i> Continue development of methodology/source data effort to simulate IP durability in laboratory and relationship to field durability.</p>								
7) Test and Evaluation (T&E) Test and Evaluation, Decontamination: Develop test and evaluation technologies and processes in support of decontamination activities. <i>FY 2009 Accomplishments:</i> Initiated and completed test and evaluation methodologies and protocols for assessing reactivity of alternative reactive material technologies and processes. Initiated and completed processes for relevant environment and relevant equipment testing for live agents and calculations for small item contact test that incorporates toxicological considerations.				1.410	0.000	0.000	0.000	0.000
Accomplishments/Planned Programs Subtotals				25.761	13.307	11.875	0.000	11.875

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 3: <i>Advanced Technology Development (ATD)</i>	R-1 ITEM NOMENCLATURE PE 0603384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ATD)</i>	PROJECT TE3: <i>TEST & EVALUATION (ATD)</i>
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C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011</u> <u>Base</u>	<u>FY 2011</u> <u>OCO</u>	<u>FY 2011</u> <u>Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To</u> <u>Complete</u>	<u>Total Cost</u>
• TE4: <i>TEST & EVALUATION (ACD&P)</i>	6.261	28.773	19.304		19.304	11.851	28.035	20.266	21.139	Continuing	Continuing
• TE5: <i>TEST & EVALUATION (SDD)</i>	37.444	36.593	15.901		15.901	12.243	4.238	14.614	15.300	Continuing	Continuing
• TE7: <i>TEST & EVALUATION (OP SYS DEV)</i>	7.037	4.870	4.813		4.813	4.779	4.750	5.660	5.615	Continuing	Continuing

D. Acquisition Strategy

N/A

E. Performance Metrics

N/A

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY			R-1 ITEM NOMENCLATURE				PROJECT				
0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 3: <i>Advanced Technology Development (ATD)</i>			PE 0603384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ATD)</i>				TR3: <i>MEDICAL RADIOLOGICAL DEFENSE (ATD)</i>				
COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
TR3: <i>MEDICAL RADIOLOGICAL DEFENSE (ATD)</i>	4.859	2.403	0.957	0.000	0.957	0.966	1.922	2.901	2.927	Continuing	Continuing

A. Mission Description and Budget Item Justification

This project (TR3) funds advanced technology development of medical countermeasures against radiological exposure. Specifically, innovative technical approaches will be used to develop, refine, and transition promising products to advanced development efforts to mitigate health consequences resulting from Acute Radiation Exposure (ARS) and Delayed Effects of Acute Radiation Exposure (DEARE). Promising products and pertinent science and technology data will be used to support Investigational New Drug (IND) applications and Food and Drug Administration (FDA) licensure processes, with an emphasis on the development of pretreatments to protect military responders in the event of a radiological incident. Research efforts and data are collaboratively shared with other government agencies so that more mature and promising product candidates will be quickly transitioned to advanced development efforts.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) Radiological Medical Countermeasures Radiation Medical Countermeasures: Develops medical countermeasures to protect the Warfighter against radiological/nuclear exposure. The Department of Defense is the only governmental agency currently developing medical prophylaxis to protect Warfighters or other responders in the event of a radiological incident. <i>FY 2009 Accomplishments:</i> Continued to evaluate at least two promising drug candidates to assess animal survival rate when exposed to lethal radiation. Evaluated efficacy of three to four therapeutic candidates and regimens that mitigate and/or treat post-radiation exposure, with emphasis on broad spectrum activity, ease of administration, and safety in non-human primates (NHPs). Continued to evaluate the preclinical efficacy and safety studies in NHPs, an assessment of drug mechanism of action,	4.859	2.371	0.957	0.000	0.957

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 3: <i>Advanced Technology Development (ATD)</i>	R-1 ITEM NOMENCLATURE PE 0603384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ATD)</i>	PROJECT TR3: <i>MEDICAL RADIOLOGICAL DEFENSE (ATD)</i>
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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p>and the determination of drug formulation according to the FDA animal rule. Evaluated promising radioprotectants and post-irradiation therapeutic agents.</p> <p><i>FY 2010 Plans:</i> Evaluate mature and promising agents for respiratory and gastrointestinal damage and repair. Demonstrate efficacy and safety in NHPs. Begin down-selection and prepare transition of one mature radioprotectant to the advanced developer, using pertinent science and technology data to support an Investigational New Drug (IND) application for eventual FDA license.</p> <p><i>FY 2011 Base Plans:</i> Continue to investigate relatively mature candidates for advanced development as medical countermeasures to prevent and treat exposure to radiation. Continue to evaluate diagnostic biodosimetry biomarkers that could be used to potentially screen mass casualties. Continue to explore the development of a biodosimetry hand-held diagnostic device that is minimally invasive, accurate, rapid, high-throughput, and suitable for medical triage. Continue development of animals models for radiation exposures useful to support FDA licensure.</p>					
2) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.	0.000	0.032	0.000	0.000	0.000
Accomplishments/Planned Programs Subtotals	4.859	2.403	0.957	0.000	0.957

C. Other Program Funding Summary (\$ in Millions)

Line Item	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	FY 2012	FY 2013	FY 2014	FY 2015	Cost To Complete	Total Cost
• MR4: <i>MEDICAL RADIOLOGICAL DEFENSE (ACD&P)</i>	4.294	0.000	0.000		0.000	0.000	0.000	0.000	0.000	Continuing	Continuing
	3.002	8.276	1.143		1.143	4.817	2.265	0.000	0.000	Continuing	Continuing

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 3: <i>Advanced Technology Development (ATD)</i>	R-1 ITEM NOMENCLATURE PE 0603384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ATD)</i>	PROJECT TR3: <i>MEDICAL RADIOLOGICAL DEFENSE (ATD)</i>
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C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011 Base</u>	<u>FY 2011 OCO</u>	<u>FY 2011 Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To Complete</u>	<u>Total Cost</u>
• MR5: <i>MEDICAL RADIOLOGICAL DEFENSE (SDD)</i>											
• TR2: <i>MEDICAL RADIOLOGICAL DEFENSE (APPLIED RESEARCH)</i>	0.525	2.897	2.884		2.884	1.904	2.855	1.913	1.903	Continuing	Continuing

D. Acquisition Strategy

N/A

E. Performance Metrics

N/A

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program								DATE: February 2010			
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 3: <i>Advanced Technology Development (ATD)</i>				R-1 ITEM NOMENCLATURE PE 0603384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ATD)</i>				PROJECT TT3: <i>TECHBASE TECHNOLOGY TRANSITION</i>			
COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
TT3: <i>TECHBASE TECHNOLOGY TRANSITION</i>	8.127	7.357	4.504	0.000	4.504	8.117	8.169	8.390	8.528	Continuing	Continuing

A. Mission Description and Budget Item Justification

This project (TT3) supports technology transition, technology experimentation and demonstration efforts, and technology readiness assessments in support of unique chemical and biological Advanced Technology Demonstrations (ATDs) and Joint Concept Technology Demonstrations (JCTDs). Within this project are two primary capability areas: 1) Experiment and Technology Demonstrations; and, 2) Technology Readiness Assessment. The Experiment and Technology Demonstrations capability area focuses on integration, testing, and assessing candidate ATDs and JCTDs and includes three thrust areas (two of which are new sub-thrust areas that consolidate legacy systems and are annotated as such below): Advanced Remediation Technologies (ART), Early Warning Military Application in Reconnaissance Systems (EW-MARS), and Comprehensive Innovative Protection (CIP). The ART addresses Chemical, Biological, and Radiological (CBR) remediation and decontamination processes and demonstrates technologies and methods to restore assets such as mobile equipment, fixed sites, critical infrastructures, personal, and equipment to operational status as a result of having reduced or eliminated CBR contamination. The EW-MARS (new thrust area) achieves enhanced command and control decision making capabilities as a result of a combined and orchestrated family of chemical and biological defense systems deployed on various platforms in remote locations. The CIP (new thrust area) transitions mature technologies to improve individual and collective protection capabilities. The Technology Readiness Assessment capability area focuses on completing manufacturing readiness assessments, technology readiness evaluations, and assessing maturity levels before transitioning ATDs and JCTDs to advanced development efforts located in Budget Activity 4 (Project TT4).

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.	0.000	0.101	0.000	0.000	0.000
2) Experiment & Technology Demonstrations	5.623	4.869	2.175	0.000	2.175

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 3: <i>Advanced Technology Development (ATD)</i>	R-1 ITEM NOMENCLATURE PE 0603384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ATD)</i>	PROJECT TT3: <i>TECHBASE TECHNOLOGY TRANSITION</i>
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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2009 Accomplishments:</i></p> <p>ART Thrust Area Continued biological decontamination technology and decision support system evaluations for IBRD. Conducted biological technology demonstrations for IBRD. Continued testing of candidate technologies for Auto Decon and HaMMER ATDs and for CBRN capability insertion into advanced development programs.</p> <p>EW Thrust Area Analyzed the capability of current- and near-term early warning technologies that may either be capable of or are required to sense CB attacks in preparation for the Early Warning/Military Applications in Reconnaissance/Surveillance ATDs.</p> <p>CIP Thrust Area Performed Design, Test and Evaluation for the IP Demo ATD.</p> <p><i>FY 2010 Plans:</i></p> <p>EW Thrust Area Conduct technology testing for EW/MARS Rapid Area Sensitive Site Reconnaissance (RASR) ATD. RASR will assess the capability to rapidly survey large areas (whole rooms, courtyards, fields) and assess and identify contamination with Chemical Warfare Agents (CWAs), Toxic Industrial Chemicals (TICs) or Non-Traditional Agents (NTAs). Conduct a technical assessment to determine if a designated WMD payload was or was not onboard a missile delivery system for the EW/MARS Post Intercept WMD Identification (PIWID) ATD.</p> <p>CIP Thrust Area Analyze the thermal burden for Warfighter protective gear in a CBRN environment as part of the CIP Low Burden Individual Protection Demonstration (IP Demo). Assessment of integrated fabric, low resistance/profile filtration, human performance prediction and assessment and low-burden air</p>					

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 3: <i>Advanced Technology Development (ATD)</i>	R-1 ITEM NOMENCLATURE PE 0603384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ATD)</i>	PROJECT TT3: <i>TECHBASE TECHNOLOGY TRANSITION</i>				
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p>purifying respirator concurrent with the Protection and Hazard Mitigation capability area (see BA2, Project CB2, Protection and Hazard Mitigation), which will support the Uniform Integrated Protective Ensemble (UIPE), and incorporate lessons into further development of integrated fabric.</p> <p><i>FY 2011 Base Plans:</i> ART Thrust Area Perform technical assessments for the ART Hazard Mitigation, Material, and Equipment Restoration (HaMMER) ATD. Incorporate results into HaMMER from testing and transition of solid oxidant and green surfactant and the Decontamination of Family Systems from the Protection and Hazard Mitigation capability area (see BA2, Project CB2, Protection and Hazard Mitigation).</p> <p>EW Thrust Area. Conduct Surety testing, technical demonstrations, and down selects for the RASR ATD.</p> <p>CIP Thrust Area Develop lessons learned from the IP Demo and inform the Protection and Hazard Mitigation capability area for future development (see BA2, Project CB2, Protection and Hazard Mitigation).</p>						
<p>3) Technology Readiness Assessment</p> <p><i>FY 2009 Accomplishments:</i> Conducted Technology Readiness Evaluations in support of remediation and restoration technology demonstrations to identify technologies in support of the ART IBRD ATD and EW MARS-JFP ATD.</p> <p><i>FY 2010 Plans:</i> Continue Technology Readiness Evaluations in support of the EW MARS-JFP ATD. For the EW RASR ATD, assess the capability to rapidly survey large areas (whole rooms, courtyards, fields) and assess and identify contamination with Chemical Warfare Agents (CWAs), Toxic Industrial Chemicals (TICs) or Non-Traditional Agents (NTAs). Build and integrate key technology components integrated</p>		2.504	2.387	2.329	0.000	2.329

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 3: <i>Advanced Technology Development (ATD)</i>	R-1 ITEM NOMENCLATURE PE 0603384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ATD)</i>	PROJECT TT3: <i>TECHBASE TECHNOLOGY TRANSITION</i>
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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
to demonstrate system level Force Protection capabilities in a Forward Operating Base scenario. Investigate the efficacy of rapid biological threat detection coupled with automatic, rapid delivery of supplies, therapeutics, and physiological monitoring equipment via unmanned systems for the CIP JMDSE ATD. <i>FY 2011 Base Plans:</i> Continue Technology Readiness Evaluations in support of the EW MARS-JFP ATD. Initiate Technology Readiness Evaluation for the CIP thrust area in preparation for a new ATD. Assess emerging innovations associated with orchestrating the response and capabilities of both individual and collective protection measures within the framework of smart networks and smart materials.					
Accomplishments/Planned Programs Subtotals	8.127	7.357	4.504	0.000	4.504

C. Other Program Funding Summary (\$ in Millions)

Line Item	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	FY 2012	FY 2013	FY 2014	FY 2015	Cost To Complete	Total Cost
• CB2: <i>CHEMICAL BIOLOGICAL DEFENSE (APPLIED RESEARCH)</i>	102.599	110.955	88.897		88.897	100.243	97.979	90.686	91.554	Continuing	Continuing
• TT4: <i>TECHBASE TECHNOLOGY TRANSITION (ACD&P)</i>	17.065	26.649	26.466		26.466	18.564	18.838	19.294	19.563	Continuing	Continuing

D. Acquisition Strategy

N/A

E. Performance Metrics

N/A

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Exhibit R-2, RDT&E Budget Item Justification: PB 2011 Chemical and Biological Defense Program										DATE: February 2010	
APPROPRIATION/BUDGET ACTIVITY				R-1 ITEM NOMENCLATURE							
0400: Research, Development, Test & Evaluation, Defense-Wide BA 4: Advanced Component Development & Prototypes (ACD&P)				PE 0603884BP: CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)							
COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
Total Program Element	69.793	209.275	277.062	0.000	277.062	182.351	198.586	169.998	177.657	Continuing	Continuing
CA4: CONTAMINATION AVOIDANCE (ACD&P)	7.703	40.186	63.347	0.000	63.347	9.093	10.754	4.742	3.978	Continuing	Continuing
CM4: HOMELAND DEFENSE (ACD&P)	0.000	0.000	9.526	0.000	9.526	0.000	0.000	0.000	0.000	Continuing	Continuing
CP4: COUNTERPROLIFERATION SUPPORT (ACD&P)	2.373	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	Continuing	Continuing
DE4: DECONTAMINATION SYSTEMS (ACD&P)	4.822	1.792	7.051	0.000	7.051	5.748	1.386	0.000	0.000	Continuing	Continuing
IP4: INDIVIDUAL PROTECTION (ACD&P)	0.000	0.000	3.172	0.000	3.172	0.000	0.000	0.000	0.000	Continuing	Continuing
IS4: INFORMATION SYSTEMS (ACD&P)	0.000	0.000	11.221	0.000	11.221	3.404	4.565	4.676	4.741	Continuing	Continuing
MB4: MEDICAL BIOLOGICAL DEFENSE (ACD&P)	7.910	102.437	136.975	0.000	136.975	130.718	131.347	115.985	113.566	Continuing	Continuing
MC4: MEDICAL CHEMICAL DEFENSE (ACD&P)	19.365	9.438	0.000	0.000	0.000	2.973	3.661	5.035	14.670	Continuing	Continuing
MR4: MEDICAL RADIOLOGICAL DEFENSE (ACD&P)	4.294	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	Continuing	Continuing
TE4: TEST & EVALUATION (ACD&P)	6.261	28.773	19.304	0.000	19.304	11.851	28.035	20.266	21.139	Continuing	Continuing
TT4: TECHBASE TECHNOLOGY TRANSITION (ACD&P)	17.065	26.649	26.466	0.000	26.466	18.564	18.838	19.294	19.563	Continuing	Continuing

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Exhibit R-2, RDT&E Budget Item Justification: PB 2011 Chemical and Biological Defense Program	DATE: February 2010
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APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>
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A. Mission Description and Budget Item Justification

Operational forces have an immediate need to survive, safely operate, and sustain operations in a Chemical and Biological (CB) agent threat environment across the continuum of global, contingency, special operations/low intensity conflict, counternarcotics, and other high risk missions. This program element supports the Advanced Component Development and Prototypes (ACD&P) of CB defensive equipment, both medical and non-medical. DoD missions for civil support operations have recently expanded and have resulted in providing focus to develop technologies to support CB counterterrorism initiatives. Projects within BA4 have been structured to consolidate Joint and Service-unique tasks within four commodity areas: contamination avoidance, force protection (individual and collective), decontamination, and medical countermeasures. ACD&P is conducted for an array of chemical/biological/toxin detection and warning systems providing early warning, collector concentrators, generic detection, improved reagents, and decontamination systems using solutions that will remove and/or detoxify contaminated material without damaging combat equipment, personnel or the environment. In the medical chemical/biological defense area, ACD&P is conducted for improved medical equipment, vaccines, and drugs essential to counteracting lethal and human performance degrading effects of chemical and biological agent threats. Specific items include improvements to nerve agent antidotes, anticonvulsants, biological agent diagnostics, and vaccines to protect against various Biological Warfare (BW) agents. This project funds development of a Transformational Rapid Drug Discovery and Development Capability (TRDDDC). Transformational Medical Technology Initiatives (TMTI) efforts in this area will include the continual build out of both a genomic sequencing and a bio-chemical informatics capability for the DoD. ACD&P also supports the Product Director Test Equipment, Strategy and Support (PD TESS) providing for the development of updated test capabilities to evaluate Chemical, Biological, Radiological and Nuclear Defense systems. Also included is the Techbase Technology Transition effort which validates high-risk/high-payoff technologies that could significantly improve warfighter capabilities. This project also funds development of candidate therapeutic medical countermeasures to mitigate the consequences of exposure to ionizing radiation due to nuclear or radiological attacks.

This Program Element focuses on efforts associated with advanced technology development used to demonstrate general military utility to include ACD&P in the areas of Non-Traditional Agents (NTA) and chemical/biological defense equipment and is correctly placed in BA4.

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Exhibit R-2, RDT&E Budget Item Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY	R-1 ITEM NOMENCLATURE
0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i>	PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>
BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	

B. Program Change Summary (\$ in Millions)

	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011 Base</u>	<u>FY 2011 OCO</u>	<u>FY 2011 Total</u>
Previous President's Budget	62.721	205.952	0.000	0.000	0.000
Current President's Budget	69.793	209.275	277.062	0.000	277.062
Total Adjustments	7.072	3.323	277.062	0.000	277.062
• Congressional General Reductions		-0.877			
• Congressional Directed Reductions		0.000			
• Congressional Rescissions	0.000	0.000			
• Congressional Adds		4.200			
• Congressional Directed Transfers		0.000			
• Reprogrammings	7.796	0.000			
• SBIR/STTR Transfer	-0.724	0.000			
• Other Adjustments	0.000	0.000	277.062	0.000	277.062

Congressional Add Details (\$ in Millions, and Includes General Reductions)

Project: CA4: CONTAMINATION AVOIDANCE (ACD&P)

Congressional Add: 1) *JBSDS Increment 2*

Congressional Add: 2) *JBTD*

Congressional Add Subtotals for Project: CA4

Project: CP4: COUNTERPROLIFERATION SUPPORT (ACD&P)

Congressional Add: 1) *JMDS*

Congressional Add Subtotals for Project: CP4

Project: DE4: DECONTAMINATION SYSTEMS (ACD&P)

Congressional Add: 1) *JMDS*

Congressional Add: 2) *JPID*

Congressional Add: 3) *JPID*

	<u>FY 2009</u>	<u>FY 2010</u>
	1.187	0.000
	0.000	0.797
	1.187	0.797
	2.373	0.000
	2.373	0.000
	1.582	0.000
	1.582	0.000
	0.000	1.792

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Exhibit R-2, RDT&E Budget Item Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>
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Congressional Add Details (\$ in Millions, and Includes General Reductions)	FY 2009	FY 2010
Congressional Add Subtotals for Project: DE4	3.164	1.792
Project: MB4: MEDICAL BIOLOGICAL DEFENSE (ACD&P)		
Congressional Add: 1) <i>Vacuum Sampling Pathogen Collection and Concentration</i>	3.164	0.000
Congressional Add: 2) <i>Broad Spectrum Therapeutic Countermeasure</i>	0.000	1.593
Congressional Add: 3) <i>CRP</i>	1.582	0.000
Congressional Add: 4) <i>TT Bio</i>	0.791	0.000
Congressional Add: 5) <i>TT Bio</i>	1.582	0.000
Congressional Add: 6) <i>TT Bio</i>	0.791	0.000
Congressional Add Subtotals for Project: MB4	7.910	1.593
Congressional Add Totals for all Projects	14.634	4.182

Change Summary Explanation

Funding: FY09 - SBIR taxes (-\$724K Various), Reprogrammings and realignments between projects (-\$791 thousand CM4; +\$2.4 million CP4; -\$3.75 thousand DE4; +\$2.373 million MB4; +\$11.304 million MC4; -\$3.740 million MR4).

FY11 - Realignment to support Decontamination Family of Systems (+\$7.1 million DE4). Inflation rate adjustment (-\$300 thousand CA4; -\$45 thousand CM4; -\$33 thousand DE4; -\$15 thousand IP4; -\$53 thousand IS4; -\$656 thousand MB4; -\$92 thousand TE4; -\$126 thousand TT4). Realignment of BA5 funding (SDD) to BA4 (ACD&P) due to change in acquisition milestone requirements (+\$57.833 million CA4; +\$9.593 million CM4; +\$3.194 million IP4; +\$11.3 million IS4). Realignment of CBDP to support test infrastructure (+\$7.7 million TE4).

Schedule: N/A

Technical: N/A

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program									DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>				R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>				PROJECT CA4: <i>CONTAMINATION AVOIDANCE (ACD&P)</i>			
COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
CA4: <i>CONTAMINATION AVOIDANCE (ACD&P)</i>	7.703	40.186	63.347	0.000	63.347	9.093	10.754	4.742	3.978	Continuing	Continuing
Quantity of RDT&E Articles	0	28	56		56	0	0	0	0		

A. Mission Description and Budget Item Justification

This Advanced Component Development and Prototypes (ACD&P) funding supports Component Advanced Development and System Integration (CAD/SI) of reconnaissance, detection, identification, and hazard prediction equipment, hardware, and software. Individual projects are: (1) Joint Biological Standoff Detector System (JBSDS), (2) Joint Biological Tactical Detection System (JBTDSD), (3) Joint Service Lightweight Standoff Chemical Agent Detector (JSLSCAD), (4) Major Defense Acquisition Program (MDAP) Support, (5) Next Generation Chemical Standoff Detection (NGCSD), and (6) Chemical Biological Radiological Nuclear Dismounted Reconnaissance Systems (CBRN DRS), formerly JNBCRS Increment 2 .

The JBSDS is employing an incremental acquisition strategy. JBSDS Increment 1 is the first standoff early warning biological detection (BD) system for the Joint Services. The system will be capable of providing near real time detection of biological attacks/incidents and standoff early detection/warning (Detect to Warn) of biological Warfare (BW) agents at fixed sites or in static mode on vehicles. It will be capable of providing standoff detection, ranging, tracking, discrimination (man-made vs. natural occurring aerosols) of BW aerosol clouds for advanced warning, reporting, and protection. The JBSDS will augment and integrate with existing BD systems to provide a BD network capable of near real time detection and warning theater-wide to limit the effects of biological agent hazards against U.S. forces at the tactical and operational levels of war. The JBSDS can be employed in support of various areas (e.g., fixed sites, Air Ports of Debarkation/Sea Ports of Debarkation (APODs/SPODs), amphibious landing sites, etc.), or on platforms (ships, aircraft or ground vehicles).

The JBSDS Increment 2 builds on the capabilities demonstrated during the development of JBSDS Increment 1. The JBSDS Increment 2 system will focus on providing 24-hour operations, improving the false alarm rate and detection sensitivity, while decreasing size, weight and power. The JBSDS Increment 2 will also integrate with the global information network to provide near real time detection and warning theater-wide to limit the effect of biological agent hazards against U.S. forces at the tactical and operational levels of war. JBSDS Increment 3 will build on Increment 2 and focus on the development of a system that will operate on-the-move on mobile platforms as determined by the Warfighter. During the Tech Development phase, JBSDS will hold competitive prototyping and key sub-system development, conduct test and evaluation of prototypes, improve agent-simulant modeling, prepare Milestone B documentation and preliminary designs.

The Joint Biological Tactical Detection System (JBTDSD) Increment 1 will develop, integrate, test and produce a lightweight, low cost biological surveillance system that will detect, collect and identify biological warfare agent aerosols. JBTDSD will provide warning through the Joint Warning And Reporting Network (JWARN) and

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<p>an archive sample for follow-on analyses. JBTDS will provide near real time local audio and visual alarm. JBTDS components will be one man portable and battery operable. JBTDS will be used organically at Brigade and below and at Forward Operating Bases (FOB) to provide notification of a hazard and enhanced battle space awareness to protect and preserve the force. When networked, JBTDS will augment existing biological detection systems to provide a theater-wide seamless array capable of biological detection, identification and warning.</p> <p>The JSLSCAD effort initiated the component improvements and the Technology Readiness Assessment (TRA) for the System of Systems (SoS) approach to address the CB early warning mission within the Next Generation Chemical Standoff Detection (NGCSD) program. The NGCSD SoS approach will increase the range of standoff detection and decrease detection time.</p> <p>The Major Defense Acquisition Program (MDAP) Support program will integrate System of Systems (SoS) solutions across the Armed Services for Major Defense Acquisition Programs (MDAP) having Chemical and Biological Radiological and Nuclear (CBRN) survivability requirements. The program will demonstrate modular, net-centric, "plug and play" capabilities for mounted and dismounted CBRN reconnaissance that will establish a common CBRN reconnaissance architecture across the services.</p> <p>The NGCSD, which was initiated under the JSLSCAD program, will provide early warning for both traditional and non-traditional chemical agent attacks at fixed sites, forward operating bases and on Service designated vehicles and ships. This effort will develop and integrate new standoff sensor technologies for future standoff systems. The detection system will interoperate with the Services and Joint Command, Control, Communications, Computers, Intelligence, Surveillance and Reconnaissance (C4ISR) architectures.</p> <p>The future increments in point detection will focus on capability gaps not addressed by M4 and M4E1 JCAD. Shipboard operation, aircraft operation, unmasking tool as well as improved decontamination verification will be primary focus areas.</p> <p>The CBRN Dismounted Reconnaissance Systems (CBRN DRS) program fills a mission critical need to enhance CBRN reconnaissance platoon capabilities. The program consists of two phases. Phase I is the Dismounted Reconnaissance (DR) Sets, Kits, and Outfits (SKO) configuration which provides an immediate critical need consisting of Commercial Off-The-Shelf (COTS) and Government Off-The-Shelf (GOTS) equipment integrated into a modular, transportable container for dismounted operations. It will form the basis for Phase II, which is the Monitoring and Survey (MS) SKO. The MS SKO will feature technology insertion, the addition of net-centric capability, and tailoring to focus on the Service-specific needs, to include Non Traditional Agent (NTA) detection.</p> <p><u>B. Accomplishments/Planned Program (\$ in Millions)</u></p>		

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) CBRN DRS <i>FY 2011 Base Plans:</i> Initiate CBRN DRS program. Initiate technology development and preliminary design to include Non Traditional Agent (NTA) and CBRN Common Sensor Interface (CCSI) efforts.		0.000	0.000	1.986	0.000	1.986
2) JBSDS Increment 2 <i>FY 2010 Plans:</i> Provide strategic, tactical planning, government system engineering, program/financial management, costing, contracting, scheduling, acquisition oversight, technical support and milestone documentation. <i>FY 2011 Base Plans:</i> Continue strategic, tactical planning, government system engineering, program/financial management, costing, contracting, scheduling, acquisition oversight, technical support and milestone documentation.		0.000	4.132	5.443	0.000	5.443
3) JBSDS Increment 2 <i>FY 2010 Plans:</i> Initiate Agent Performance Assessment and Cross Section Measurements. <i>FY 2011 Base Plans:</i> Continue Agent Performance Assessment and Cross Section Measurements.		0.000	1.250	1.500	0.000	1.500
4) JBSDS Increment 2 <i>FY 2010 Plans:</i> Initiate Increment 2 Modeling and Simulation in support of Agent Performance Assessment. <i>FY 2011 Base Plans:</i> Continuing Increment 2 Modeling and Simulation in support of Agent Performance Assessment.		0.000	1.500	2.000	0.000	2.000

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
5) JBSDS Increment 2 <i>FY 2010 Plans:</i> Initiate Technology Development and Preliminary Designs including competitive prototyping. <i>FY 2011 Base Plans:</i> Continue Technology Development and Preliminary Designs and competitive prototyping.		0.000	9.701	10.336	0.000	10.336
6) JBSDS Increment 2 <i>FY 2010 Plans:</i> Provide developmental test organizations funding to support test planning and test support for evaluation of the competitive prototypes.		0.000	1.100	0.000	0.000	0.000
7) JBSDS Increment 2 <i>FY 2011 Base Plans:</i> Initiate pre-Milestone B contractual activities including development of proposal package, release of draft request for proposal (RFP), update RFP with CDD requirements and comments from industry, conduct government evaluation team source selection training, and system engineering activities leading up to Engineering and Manufacturing Development phase.		0.000	0.000	2.425	0.000	2.425
8) JBSDS Increment 2 <i>FY 2011 Base Plans:</i> Initiate validation of simulants, models, and test support equipment including referee equipment development for the evaluation of competitive prototypes and advance development hardware.		0.000	0.000	6.447	0.000	6.447
9) JBTDS		2.129	3.651	3.698	0.000	3.698

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2009 Accomplishments:</i> Continued to provide strategic/tactical planning, government systems engineering, program/financial management, costing, technology assessment, contracting, scheduling, acquisition oversight and technical support.</p> <p><i>FY 2010 Plans:</i> Continue to provide strategic/tactical planning, government systems engineering, program/financial management, costing, technology assessment, contracting, scheduling, acquisition oversight and technical support.</p> <p><i>FY 2011 Base Plans:</i> Continue to provide strategic/tactical planning, government systems engineering, program/financial management, costing, technology assessment, contracting, scheduling, acquisition oversight and technical support.</p>						
10) JBTDS <i>FY 2009 Accomplishments:</i> Conducted Technology Readiness Evaluations on identifier candidates.		1.045	0.000	0.000	0.000	0.000
11) JBTDS <i>FY 2009 Accomplishments:</i> Continued Pre-Milestone A activities, analysis and risk reduction demonstrations. <i>FY 2010 Plans:</i> Complete Pre-Milestone A activities, analysis and risk reduction demonstrations.		1.500	4.007	0.000	0.000	0.000
12) JBTDS		0.235	0.000	0.000	0.000	0.000

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2009 Accomplishments:</i> Continued network algorithm development/demonstration.						
13) JBTDS <i>FY 2011 Base Plans:</i> Initiate government testing and analysis, to include Manufacturing Readiness Assessment and Technology Readiness Assessment of hardware submitted for evaluation in Competitive Prototyping.		0.000	0.000	2.700	0.000	2.700
14) JBTDS <i>FY 2010 Plans:</i> Initiate competitive prototyping contracts effort for JBTDS Increment 1 prototypes. <i>FY 2011 Base Plans:</i> Continue competitive prototyping contracts effort for JBTDS Increment 1 prototypes.		0.000	1.800	1.500	0.000	1.500
15) JBTDS <i>FY 2011 Base Plans:</i> Initiate technology development phase activities to mature technologies that are at Technology Readiness Level 4 (TRL 4).		0.000	0.000	0.900	0.000	0.900
16) JBTDS <i>FY 2010 Plans:</i> Conduct field demonstration analysis for end to end integrated candidates at government test ranges.		0.000	1.106	0.000	0.000	0.000
17) JBTDS		0.000	0.000	1.434	0.000	1.434

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2011 Base Plans:</i> Initiate activities to support Milestone B Engineering and Manufacturing Development (EMD) phase requirements and milestone documentation development.						
18) JBTDS <i>FY 2011 Base Plans:</i> Initiate assay transition with Critical Reagents Program (CRP) for identifier to include emerging bio-surveillance threats approved by threat agencies.		0.000	0.000	2.500	0.000	2.500
19) JBTDS <i>FY 2011 Base Plans:</i> Initiate pre-MS B contractual efforts including development of proposal package, release draft Request For Proposal (RFP), prepare final EMD RFP, release RFP, conduct source selection training, conduct source selection and complete proposal evaluations.		0.000	0.000	3.400	0.000	3.400
20) JCAD <i>FY 2011 Base Plans:</i> Initiate Analysis of Alternatives for the future increments in point detection.		0.000	0.000	1.986	0.000	1.986
21) JSLSCAD <i>FY 2009 Accomplishments:</i> Conducted strategic/tactical planning, systems engineering, and technology assessment for Next Generation Chemical Standoff Detection (NGCSD).		0.907	0.000	0.000	0.000	0.000
22) JSLSCAD		0.577	0.000	0.000	0.000	0.000

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2009 Accomplishments:</i> Conducted Testing Evaluation and Methodology Analysis for NGCSD.						
23) JSLSCAD <i>FY 2009 Accomplishments:</i> Integrated Requirements Analysis, Development, and Demonstration for NGCSD.		0.123	0.000	0.000	0.000	0.000
24) MDAP SPRT Catalytic Oxidation (CatOx) Technology Demonstration of improved air purification for the Abrams Main Battle Tank. <i>FY 2010 Plans:</i> Initiate transition of CatOx air purification process for design to increase CBRN defensive capability and reduce logistical costs. <i>FY 2011 Base Plans:</i> Initiate developmental (shock, vibration, temperature, dust environment) testing of CatOx prototypes in the Abrams Main Battle Tank operating environment.		0.000	1.100	0.900	0.000	0.900
25) MDAP SPRT Chemical, Biological, Radiological and Nuclear (CBRN) Capabilities Analysis <i>FY 2010 Plans:</i> Conduct CBRN Capabilities Analysis for Ground Combat Vehicle. <i>FY 2011 Base Plans:</i> Conduct CBRN Capabilities Analysis for the Ship to Shore Connector and Ground Combat Vehicle.		0.000	0.200	0.500	0.000	0.500

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
26) MDAP SPRT Chemical, Biological, Radiological and Nuclear (CBRN) Material Solutions Analysis. <i>FY 2010 Plans:</i> Conduct CBRN Material Solutions Analysis for Ground Combat Vehicle. <i>FY 2011 Base Plans:</i> Complete CBRN Material Solutions Analysis for Ground Combat Vehicle. Initiate the CBRN Material Solutions Analysis for Ship to Shore Connector, Apache Block Three, and Next Generation Cruiser (CG(X)).		0.000	0.216	1.577	0.000	1.577
27) NGCSD <i>FY 2010 Plans:</i> Initiate sensor prototype development. <i>FY 2011 Base Plans:</i> Continue sensor prototype development.		0.000	4.500	6.015	0.000	6.015
28) NGCSD <i>FY 2010 Plans:</i> Conduct Technology Readiness Evaluation (TRE) and support of Analysis of Alternatives (AoA). <i>FY 2011 Base Plans:</i> Continue support of Analysis of Alternatives (AoA).		0.000	1.833	0.300	0.000	0.300
29) NGCSD		0.000	2.800	1.500	0.000	1.500

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2010 Plans:</i> Initiate engineering support.						
<i>FY 2011 Base Plans:</i> Continue engineering support.						
30) NGCSD		0.000	0.000	2.700	0.000	2.700
<i>FY 2011 Base Plans:</i> Initiate Pre-MS B prototype procurement (3 each of 3 technologies at a cost of \$300 thousand each).						
31) NGCSD		0.000	0.000	0.400	0.000	0.400
<i>FY 2011 Base Plans:</i> Initiate planning for Early Operational Assessment (EOA).						
32) NGCSD		0.000	0.000	1.200	0.000	1.200
<i>FY 2011 Base Plans:</i> Initiate prototype Developmental Test (DT) planning and execution.						
33) SBIR		0.000	0.493	0.000	0.000	0.000
<i>FY 2010 Plans:</i> Small Business Innovative Research.						
Accomplishments/Planned Programs Subtotals		6.516	39.389	63.347	0.000	63.347
		FY 2009	FY 2010			
Congressional Add: 1) JBSDS Increment 2		1.187	0.000			

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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010
<i>FY 2009 Accomplishments:</i> Initiated Testing and Support Equipment Development.		
Congressional Add: 2) JBTDSD <i>FY 2010 Plans:</i> Initiate Sample Preparation for Biological Detection	0.000	0.797
Congressional Adds Subtotals	1.187	0.797

C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011</u> <u>Base</u>	<u>FY 2011</u> <u>OCO</u>	<u>FY 2011</u> <u>Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To</u> <u>Complete</u>	<u>Total Cost</u>
• CA5: <i>CONTAMINATION AVOIDANCE (SDD)</i>	46.316	78.042	124.936		124.936	117.729	110.250	89.493	58.830	Continuing	Continuing
• JC0100: <i>JOINT BIO POINT DETECTION SYSTEM (JBPDS)</i>	75.545	41.976	43.555		43.555	41.252	52.776	73.164	71.894	Continuing	Continuing
• JC0101: <i>JS CHEM/BIO/RAD AGENT WATER MONITOR (JCBRAWM)</i>	6.000	3.184	0.000		0.000	0.000	0.000	0.000	0.000	Continuing	Continuing
• JC0250: <i>JOINT BIO STANDOFF DETECTOR SYSTEM (JBSDS)</i>	4.000	0.000	0.000		0.000	0.273	19.840	20.834	35.728	Continuing	Continuing
• JC4500: <i>NEXT GENERATION CHEMICAL STANDOFF DETECTION (NGCSD)</i>	0.000	0.000	0.000		0.000	0.000	9.840	12.120	21.799	Continuing	Continuing
• JF0100: <i>JOINT CHEMICAL AGENT DETECTOR (JCAD)</i>	58.406	27.694	40.071		40.071	45.805	52.762	53.330	63.217	Continuing	Continuing
• JN0900: <i>NON TRADITIONAL AGENT DETECTION (NTAD)</i>	0.000	0.000	4.178		4.178	4.075	3.376	6.745	8.918	Continuing	Continuing

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C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011</u> <u>Base</u>	<u>FY 2011</u> <u>OCO</u>	<u>FY 2011</u> <u>Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To</u> <u>Complete</u>	<u>Total Cost</u>
• MC0100: <i>JOINT NBC RECONNAISSANCE SYSTEM (JNBCRS)</i>	32.699	32.421	22.511		22.511	65.779	122.214	50.385	0.000	Continuing	Continuing
• MC0101: <i>CBRN DISMOUNTED RECONNAISSANCE SYSTEMS (CBRN DRS)</i>	0.000	11.415	15.414		15.414	24.056	33.504	56.718	53.938	Continuing	Continuing
• MC0102: <i>JOINT CONTAMINATED SURFACE DETECTOR (JCSD)</i>	0.000	0.000	0.000		0.000	0.000	0.000	5.288	40.072	Continuing	Continuing
• MX0001: <i>JOINT BIO TACTICAL DETECTION SYSTEM (JBTDS)</i>	0.000	0.000	0.000		0.000	8.080	19.060	29.237	33.933	Continuing	Continuing

D. Acquisition Strategy

CBRN DRS

The Chemical Biological Radiological Nuclear Dismounted Reconnaissance Systems (CBRN DRS) program uses spiral development with an evolutionary component/suite upgrade acquisition approach. Funding finalizes the design to provide the Services with enhanced full spectrum CBRN detection capability to support strategic, operational, and tactical objectives at lower life cycle costs. CBRN DRS will enhance the Situational Awareness (SA) by providing a dismounted ability to detect chemical, biological and radiological hazards across the Range of Military Operations (ROMO) and employ contamination avoidance activities to prevent disruption to operations and organizations.

JBSDS

The Joint Bio Stand-off Detector System (JBSDS) is employing an incremental acquisition strategy. JBSDS Increment 1 is the first standoff early warning biological detection (BD) system for the Joint Services. The JBSDS Increment 2 system will focus on providing 24-hour operations (Increment 1 is night-time only), improving the false alarm rate and detection sensitivity, while decreasing size, weight and power. The JBSDS Increment 2 will also integrate with the global information network to provide near real time detection and warning theater-wide to limit the effect of biological agent hazards against U.S. forces at the tactical and operational levels of war. JBSDS Increment 3 will build on Increment 2 and focus on the development of a system that will operate on-the-move on mobile platforms as determined by the Warfighter.

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<p>JBTDS</p> <p>The Joint Biological Tactical Detection (JBTDS) program will pursue an evolutionary incremental approach to provide capability to the Warfighter. The JBTDS program will develop, integrate, test, procure and field systems that improve biological aerosol detection, identification, and sampling capabilities. The JBTDS program will also reduce size, weight, power consumption, and logistic footprint over current systems. Test Readiness Evaluations (TRE) and Competitive Prototyping will support the JBTDS Engineering and Manufacturing Development (EMD) phase by identifying mature technologies and reducing overall risk. Modeling and simulation tools will be used in order to lower program risks, reduce costs and ensure a higher confidence in selected technologies.</p> <p>JCAD</p> <p>The current strategy employs a product improvement of the M4 JCAD to reduce Life Cycle costs, transition to a competitive procurement contract, and attain objective capability. Three competitive fixed-price contracts for the M4E1 were awarded in Sep 2007 for prototypes and options for full rate production. Competitive prototype testing was conducted and one system was selected for continued development. The production options will be exercised in FY11 following a successful production cut-in decision.</p> <p>JSLSCAD</p> <p>The acquisition strategy for the Joint Service Lightweight Standoff Chemical Agent Detector (JSLSCAD) production phase focused upon a dual path to procure required systems and concurrently develop and test system improvements to increase the military utility. The Milestone Decision Authority (MDA) approved procurement of additional JSLSCAD LRIP systems in February 2008. The Government awarded a Fixed Price Incentive contract to GD-ATP in July 2008 for production of systems to fulfill the Nuclear Biological Chemical Reconnaissance Vehicle (NBCRV) Extended LRIP requirements and additional delivery orders will be exercised for full rate production of systems to fulfill the remaining NBCRV requirements. The JSLSCAD program office awarded multiple contracts to support system engineering, software development, test and evaluation, and system support efforts to increase standoff detection capabilities to rapidly respond to evolving system integration requirements with minimal contractual lead time. All these efforts are being integrated into the Next Generation Chemical Standoff Detection (NGCSD) program.</p> <p>MDAP SPRT</p> <p>Major Defense Acquisition Program (MDAP) Support program will integrate System of Systems (SoS) solutions across the Armed Service's for Major Defense Acquisition Programs (MDAP) having Chemical and Biological Radiological and Nuclear (CBRN) survivability requirements. The MDAP program will achieve these SoS solutions by: (1) leading CBRN architecture development and System Engineering efforts that result in enterprise concepts that address requirements; (2) establishing agreements with the MDAPs on roles and responsibilities with respect to funding deliverables and integration; (3) demonstrating modular, net-centric,</p>		

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT CA4: <i>CONTAMINATION AVOIDANCE (ACD&P)</i>
<p>"plug and play" capabilities for mounted and dismounted CBRN reconnaissance requirements; (4) developing design and test schedules which synchronize support for CBRN capability integration with MDAPs' schedules; and (5) providing integrated program management across the CBRN commodity areas to deliver capabilities on time that support MDAP goals.</p> <p>NGCSD</p> <p>The Next Generation Chemical Standoff Detection (NGCSD) program, which was initiated under the JSLSCAD program, will award Indefinite Delivery/Indefinite Quantity contract(s) to support system engineering, software development, test and evaluation, and system support efforts to increase standoff detection capabilities. This contract type will allow the program office to rapidly respond to evolving system integration requirements and emerging test results with minimal contractual lead time. This will optimize the program goal of inserting the latest software and standoff detection technology into the host platforms in the shortest possible time.</p> <p><u>E. Performance Metrics</u> N/A</p>		

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT CA4: <i>CONTAMINATION AVOIDANCE (ACD&P)</i>
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Product Development (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** CBRN DRS - HW C - CBRN DRS Preliminary Design	C/FP	TBD	0.000	0.000		1.386	Jan 2011	0.000		1.386	0.000	1.386	0.000
** JBSDS - HW SB - Technology Development and Preliminary Designs	C/FPI	TBD	0.000	9.701	Jan 2010	10.336	Oct 2010	0.000		10.336	0.000	20.037	0.000
** JBTDS - HW S - Competitive Prototype Contract	C/FFP	TBD	0.000	1.800	Jul 2010	1.500	Oct 2010	0.000		1.500	0.000	3.300	0.000
HW S - EMD Contract development and Source Selection Activities	C/CPIF	TBD	0.000	0.000		3.400	Apr 2011	0.000		3.400	0.000	3.400	0.000
HW S - Tech Development	C/FP	TBD	0.000	0.000		0.900	Jan 2011	0.000		0.900	0.000	0.900	0.000
HW S - Automated Sample Prep	SS/FFP	TBD	0.000	0.797	Jul 2010	0.000		0.000		0.000	0.000	0.797	0.000
** JCAD - SW SB - Initiate Sensor Prototype Development	C/CPFF	TBD	0.000	0.000		0.750	Jan 2011	0.000		0.750	0.000	0.750	0.000
** MDAP SPRT - HW S - Catalytic Oxidation (CatOx) Technology Demonstration	C/CPFF	Honeywell Corporation Phoenix, AZ	0.000	1.100	Jan 2010	0.900	Jan 2011	0.000		0.900	0.000	2.000	0.000
** NGCSD - HW SB - Initiate Sensor Prototype Development	C/CPFF	TBD	0.000	4.500	Apr 2010	6.015	Apr 2011	0.000		6.015	0.000	10.515	0.000

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT CA4: <i>CONTAMINATION AVOIDANCE (ACD&P)</i>
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Product Development (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
SW SB - Pre-MS B Prototype Acquisition (3 each of 3 technologies)	C/CPFF	TBD	0.000	0.000		2.700	Jan 2011	0.000		2.700	0.000	2.700	0.000
Subtotal			0.000	17.898		27.887		0.000		27.887	0.000	45.785	0.000

Remarks

Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** JBSDS - ES S - INC 2 - Modeling & Simulation Test Support	C/CPFF	Bricks Sigal & Miller Inc., Kennett Square	0.000	0.400	Jan 2010	0.500	Jan 2011	0.000		0.500	0.000	0.900	0.000
ES S - INC 2 - Modeling & Simulation Test Support	C/CPFF	NAVSEA Johns Hopkins Applied Physics Lab, Baltimore	0.000	1.500	Jan 2010	1.436	Jan 2011	0.000		1.436	0.000	2.936	0.000
ES S - INC 2 - Modeling & Simulation Test Support #2	MIPR	Sandia National Lab Albuquerque, NM	0.000	0.500	Jan 2010	0.936	Jan 2011	0.000		0.936	0.000	1.436	0.000
ES S - INC 2 - Modeling, Simulation & Data Analysis	MIPR	Various	0.000	0.350	Oct 2009	0.000		0.000		0.000	0.000	0.350	0.000

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT CA4: <i>CONTAMINATION AVOIDANCE (ACD&P)</i>
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Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** JBTD S - ES S - Pre MS A Analysis & Activities	MIPR	Various	1.714	3.229	Jan 2010	0.000		0.000		0.000	0.000	4.943	0.000
ES S - EMD Activities & Analysis and Document Development	MIPR	Various	0.000	0.000		1.434	Jan 2011	0.000		1.434	0.000	1.434	0.000
ES S - Assay Development	MIPR	CRP Frederick, MD	0.000	0.000		2.500	Apr 2011	0.000		2.500	0.000	2.500	0.000
ES S - MS A Document Development	C/FFP	TBD	0.000	0.778	Jan 2010	0.000		0.000		0.000	0.000	0.778	0.000
** JCAD - TD/D SB - MS A	MIPR	TBD	0.000	0.000		0.500	Oct 2010	0.000		0.500	0.000	0.500	0.000
** MDAP SPRT - ES S - CBRN Capability Analysis	MIPR	Various	0.000	0.200	Jan 2010	0.500	Jan 2011	0.000		0.500	0.000	0.700	0.000
ES S - CBRN Material Solutions Analysis	MIPR	Various	0.000	0.216	Jan 2010	1.577	Jan 2011	0.000		1.577	0.000	1.793	0.000
** NGCSD - TD/D S - AoA	MIPR	TBD	0.000	0.500	Jan 2010	0.300	Jan 2011	0.000		0.300	0.000	0.800	0.000
Subtotal			1.714	7.673		9.683		0.000		9.683	0.000	19.070	0.000

Remarks

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT CA4: <i>CONTAMINATION AVOIDANCE (ACD&P)</i>
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Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** JBSDS - OTHT SB - INC 2 - Developmental Testing Support	MIPR	DPG Dugway, UT	0.000	1.100	Jan 2010	5.000	Jan 2011	0.000		5.000	0.000	6.100	0.000
OTHT SB - Networking algorithm development	MIPR	MA Institute of Technology - Lincoln Labs Boston, MA	0.000	0.000		0.500	Jan 2011	0.000		0.500	0.000	0.500	0.000
OTHT SB - Requirements Testing	MIPR	WSMR Nevada Test Site, NV	0.000	0.000		1.000	Jan 2011	0.000		1.000	0.000	1.000	0.000
OTHT SB - Agent performance analysis support	MIPR	DPG Dugway, UT	0.000	0.000		1.000	Jan 2011	0.000		1.000	0.000	1.000	0.000
OTHT SB - Agent performance analysis	MIPR	Johns Hopkins - Applied Physics Lab Baltimore, MD	0.000	0.000		2.000	Jan 2011	0.000		2.000	0.000	2.000	0.000
** JBTDS - OTHT SB - Competitive Prototype Testing	MIPR	DPG Dugway, UT	0.000	0.000		2.700	Oct 2010	0.000		2.700	0.000	2.700	0.000
DTE SB - Field Demonstration	MIPR	DPG Dugway, UT	0.000	1.106	Jan 2010	0.000		0.000		0.000	0.000	1.106	0.000
** JCAD - OTHT SB - Technology Demo	MIPR	Various	0.000	0.000		0.500	Apr 2011	0.000		0.500	0.000	0.500	0.000
** NGCSD - OTHT SB - Conduct TRE	MIPR	Various	0.000	1.333	Apr 2010	0.000		0.000		0.000	0.000	1.333	0.000

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT CA4: <i>CONTAMINATION AVOIDANCE (ACD&P)</i>
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Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
OTHT S - Conduct EOA	MIPR	Various	0.000	0.000		0.400	Jan 2011	0.000		0.400	0.000	0.400	0.000
DTE S - DT Planning and Execution	MIPR	Various	0.000	0.000		1.200	Jan 2011	0.000		1.200	0.000	1.200	0.000
Subtotal			0.000	3.539		14.300		0.000		14.300	0.000	17.839	0.000

Remarks

Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** CBRN DRS - PM/MS S - JPM CA	MIPR	JPM CA APG, MD	0.000	0.000		0.600	Oct 2010	0.000		0.600	0.000	0.600	0.000
** JBSDS - PM/MS S - JPM BD	MIPR	JPM BD APG, MD	0.000	1.600	Jan 2010	2.207	Jan 2011	0.000		2.207	0.000	3.807	0.000
PM/MS S - PM/MS Other Government Agencies	MIPR	USN USMC, USAF	0.000	1.632	Jan 2010	0.548	Jan 2011	0.000		0.548	0.000	2.180	0.000
PM/MS S - JPEO Management Support	Allot	JPEO Falls Church, VA	0.000	0.900	Jul 2010	2.688	Jul 2011	0.000		2.688	0.000	3.588	0.000
** JBTDS - PM/MS S - JPM BD, APG, MD	MIPR	JPM BD APG, MD	3.059	3.651	Jan 2010	3.698	Oct 2010	0.000		3.698	0.000	10.408	0.000

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** JCAD - PM/ MS SB - Program Management and Systems Engineering Support	MIPR	JPM NBC CA APG, MD	0.000	0.000		0.236	Oct 2010	0.000		0.236	0.000	0.236	0.000
** NGCSD - PM/ MS S - Program Management and Systems Engineering Support	MIPR	JPM NBC CA APG, MD	0.000	2.800	Oct 2009	1.500	Oct 2010	0.000		1.500	0.000	4.300	0.000
** ZSBIR - SBIR/STTR - Aggregated from ZSBIR-SBIR/STTR	MIPR	HQ AMC, Alexandria	0.000	0.493		0.000		0.000		0.000	0.000	0.493	0.000
Subtotal			3.059	11.076		11.477		0.000		11.477	0.000	25.612	0.000

Remarks

	Total Prior Years Cost	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
Project Cost Totals	4.773	40.186	63.347	0.000	63.347	0.000	108.306	0.000

Remarks

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT CA4: <i>CONTAMINATION AVOIDANCE (ACD&P)</i>
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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
** CBRN DRS - Monitoring and Survey (MS) Material Development Decision											■																	
Monitoring and Survey (MS) Milestone B																												
** JBSDS - Increment 2 - Technology Modeling	■	■	■	■	■	■	■																					
Increment 2 - Pre-Milestone A	■	■	■	■	■	■	■																					
Increment 2 - Milestone A							■																					
Increment 2 - Technology Development							■	■	■	■	■	■	■	■	■													
Increment 2 - Preliminary Design Review																■												
Increment 2 - Milestone B																	■											
Increment 2 - Engineering & Manufacturing Development																		■	■	■	■	■	■	■	■	■	■	
Increment 2 - Milestone C																											■	
Increment 2 - LRIP																											■	
** JBTDS - Materiel Development Decision				■																								
MS A Decision							■																					
Competitive Prototyping Contract Award									■																			
Competitive Prototyping Testing											■	■																
Capability Development Document												■																
PDR													■															
MS B Decision																											■	

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT CA4: <i>CONTAMINATION AVOIDANCE (ACD&P)</i>
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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
EMD Contract Award											■																	
Developmental Testing											■	■	■	■	■	■	■	■	■									
CDR															■													
MS C Decision																			■									
MS C Contract Award																			■									
Production Verification Test																				■								
IOT&E																				■	■	■						
FRP Decision																					■							
FRP Contract Award																						■						
IOC																										■		
** JCAD - Future Generation Chemical Point Detection - Materiel Development Decision (MDD)											■																	
Future Generation Chemical Point Detection - MS A											■																	
Future Generation Chemical Point Detection - Prototype Development and Demo											■	■																
** JSLSCAD - SoS Technology Demo	■																											
SoS Program	■	■	■	■																								
SoS Operational Demo			■	■																								
** MDAP SPRT - Catox Tech Demonstration for Abrams Main Battle Tank					■	■	■	■	■	■	■	■	■															

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT CA4: <i>CONTAMINATION AVOIDANCE (ACD&P)</i>
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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
Advance Component Prototype Development of JSF Decontamination				■	■	■	■	■	■	■	■	■	■															
** NGCSD - Material Development Decision (MDD)						■																						
Analysis of Alternatives (AoA)							■	■																				
MS A										■																		
Competitive Prototyping										■	■	■	■															
Preliminary Design Review (PDR)																■												
MS B																■												

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT CA4: <i>CONTAMINATION AVOIDANCE (ACD&P)</i>
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Schedule Details

Event	Start		End	
	Quarter	Year	Quarter	Year
** CBRN DRS - Monitoring and Survey (MS) Material Development Decision	2	2011	2	2011
Monitoring and Survey (MS) Milestone B	4	2012	4	2012
** JBSDS - Increment 2 - Technology Modeling	4	2004	3	2010
Increment 2 - Pre-Milestone A	1	2008	3	2010
Increment 2 - Milestone A	3	2010	3	2010
Increment 2 - Technology Development	3	2010	3	2012
Increment 2 - Preliminary Design Review	2	2012	2	2012
Increment 2 - Milestone B	3	2012	3	2012
Increment 2 - Engineering & Manufacturing Development	3	2012	1	2015
Increment 2 - Milestone C	1	2015	1	2015
Increment 2 - LRIP	1	2015	2	2017
** JBTDS - Materiel Development Decision	4	2009	4	2009
MS A Decision	3	2010	3	2010
Competitive Prototyping Contract Award	4	2010	4	2010
Competitive Prototyping Testing	1	2011	2	2011
Capability Development Document	1	2011	1	2011
PDR	2	2011	2	2011
MS B Decision	3	2011	3	2011

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT CA4: <i>CONTAMINATION AVOIDANCE (ACD&P)</i>
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Event	Start		End	
	Quarter	Year	Quarter	Year
EMD Contract Award	3	2011	3	2011
Developmental Testing	3	2011	3	2013
CDR	4	2012	4	2012
MS C Decision	3	2013	3	2013
MS C Contract Award	3	2013	3	2013
Production Verification Test	1	2014	1	2014
IOT&E	1	2014	3	2014
FRP Decision	3	2014	3	2014
FRP Contract Award	3	2014	3	2014
IOC	3	2015	3	2015
** JCAD - Future Generation Chemical Point Detection - Materiel Development Decision (MDD)	1	2011	1	2011
Future Generation Chemical Point Detection - MS A	1	2011	1	2011
Future Generation Chemical Point Detection - Prototype Development and Demo	2	2011	3	2011
** JSLSCAD - SoS Technology Demo	4	2008	1	2009
SoS Program	4	2008	4	2009
SoS Operational Demo	3	2009	4	2009
** MDAP SPRT - Catox Tech Demonstration for Abrams Main Battle Tank	1	2010	4	2011
Advance Component Prototype Development of JSF Decontamination	4	2009	1	2012
** NGCSD - Material Development Decision (MDD)	2	2010	2	2010
Analysis of Alternatives (AoA)	3	2010	4	2010

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT CA4: <i>CONTAMINATION AVOIDANCE (ACD&P)</i>
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Event	Start		End	
	Quarter	Year	Quarter	Year
MS A	1	2011	1	2011
Competitive Prototyping	1	2011	4	2011
Preliminary Design Review (PDR)	4	2011	4	2011
MS B	1	2012	1	2012

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT CM4: <i>HOMELAND DEFENSE (ACD&P)</i>
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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
CM4: <i>HOMELAND DEFENSE (ACD&P)</i>	0.000	0.000	9.526	0.000	9.526	0.000	0.000	0.000	0.000	Continuing	Continuing
Quantity of RDT&E Articles											

A. Mission Description and Budget Item Justification

This project funds initial development of the Common Analytical Laboratory System (CALs) to include evaluation and selection of subsystems (analytical detection, laboratory information management, data fusion, engineering controls) as well as development of a set of modular designed configurations for system level prototyping utilizing open system architecture. In addition, it provides for validation and demonstration of desired functional capabilities.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) CALS <i>FY 2011 Base Plans:</i> Continue program office planning and programming.	0.000	0.000	0.513	0.000	0.513
2) CALS <i>FY 2011 Base Plans:</i> Continue system engineering and logistics support.	0.000	0.000	1.017	0.000	1.017
3) CALS <i>FY 2011 Base Plans:</i> Complete subsystem design and development - open architecture design analytics and laboratory information management.	0.000	0.000	1.837	0.000	1.837
4) CALS	0.000	0.000	5.511	0.000	5.511

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT CM4: <i>HOMELAND DEFENSE (ACD&P)</i>
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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2011 Base Plans:</i> Develop system prototypes - alternative system configuration approaches.					
5) CALS <i>FY 2011 Base Plans:</i> Initiate system demonstration and validation.	0.000	0.000	0.648	0.000	0.648
Accomplishments/Planned Programs Subtotals	0.000	0.000	9.526	0.000	9.526

C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011 Base</u>	<u>FY 2011 OCO</u>	<u>FY 2011 Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To Complete</u>	<u>Total Cost</u>
• CM5: <i>HOMELAND DEFENSE (SDD)</i>	3.897	8.638	1.166		1.166	3.822	0.000	2.361	2.413	Continuing	Continuing
• JS0004: <i>WMD - CIVIL SUPPORT TEAMS (WMD CST)</i>	8.300	11.765	39.862		39.862	33.402	37.398	44.817	47.159	Continuing	Continuing
• JS0500: <i>CB INSTALLATION/ FORCE PROTECTION PROGRAM (FORCE PROT)</i>	80.103	53.623	50.773		50.773	60.324	59.836	57.840	54.455	Continuing	Continuing

D. Acquisition Strategy

CALS

The Common Analytical Laboratory System (CALs) will follow an incremental approach designed to address known joint force capability requirements for Chemical, Biological, Radiological and Nuclear (CBRN) detection which includes Toxic Industrial Chemicals (TICs), Toxic Industrial Materials (TIMs), Chemical Warfare Agents (CWAs), Biological Warfare Agents (BWAs). CALs will address situational awareness by leveraging efforts underway with JPEO-CBD to the extent possible. CALs will accommodate these component requirements within a modular and scalable concept framework.

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
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E. Performance Metrics

N/A

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT CM4: <i>HOMELAND DEFENSE (ACD&P)</i>
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Product Development (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** CALS - HW SB - Analytical Subsystem Evaluation and Selection	C/CPIF	TBD	0.000	0.000		0.473	Jan 2011	0.000		0.473	0.000	0.473	0.000
SW SB - Information Technology Subsystem Evaluation and Selection	C/CPIF	TBD	0.000	0.000		0.480	Jan 2011	0.000		0.480	0.000	0.480	0.000
HW S - Analytical Protocol Development	MIPR	TBD	0.000	0.000		0.884	Jan 2011	0.000		0.884	0.000	0.884	0.000
HW S - CALS Prototype Systems	C/CPIF	TBD	0.000	0.000		5.511	Jan 2011	0.000		5.511	0.000	5.511	0.000
Subtotal			0.000	0.000		7.348		0.000		7.348	0.000	7.348	0.000

Remarks

Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** CALS - ES S - Engineering Support System - CALS	MIPR	TBD	0.000	0.000		0.513	Oct 2010	0.000		0.513	0.000	0.513	0.000
Subtotal			0.000	0.000		0.513		0.000		0.513	0.000	0.513	0.000

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT CM4: <i>HOMELAND DEFENSE (ACD&P)</i>
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Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			

Remarks

Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** CALS - OTHT S - CALS Prototype Demonstration and Validation	MIPR	TBD	0.000	0.000		0.648	Apr 2011	0.000		0.648	0.000	0.648	0.000
Subtotal			0.000	0.000		0.648		0.000		0.648	0.000	0.648	0.000

Remarks

Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** CALS - PM/MS S - Program Office	MIPR	TBD	0.000	0.000		1.017	Jan 2011	0.000		1.017	0.000	1.017	0.000

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT CM4: <i>HOMELAND DEFENSE (ACD&P)</i>
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Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
- Planning and Programming													
Subtotal			0.000	0.000		1.017		0.000		1.017	0.000	1.017	0.000

Remarks

Project Cost Totals	Total Prior Years Cost	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
		0.000	0.000	9.526	0.000	9.526	0.000	9.526

Remarks

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT CM4: <i>HOMELAND DEFENSE (ACD&P)</i>
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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
** CALS - CALS Design, Development and Integration					■	■	■	■	■	■	■	■	■	■														
CALS System Demonstration														■														
CALS MDD					■																							
CALS Milestone A							■																					
CALS Milestone C														■														

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT CM4: <i>HOMELAND DEFENSE (ACD&P)</i>
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Schedule Details

Event	Start		End	
	Quarter	Year	Quarter	Year
** CALS - CALS Design, Development and Integration	1	2010	2	2012
CALS System Demonstration	2	2012	2	2012
CALS MDD	1	2010	1	2010
CALS Milestone A	3	2010	3	2010
CALS Milestone C	2	2012	2	2012

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT CP4: <i>COUNTERPROLIFERATION SUPPORT (ACD&P)</i>
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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
CP4: <i>COUNTERPROLIFERATION SUPPORT (ACD&P)</i>	2.373	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	Continuing	Continuing
Quantity of RDT&E Articles											

A. Mission Description and Budget Item Justification

This project provides for the Joint Material Decontamination System (JMDS) as a Congressional Interest Item in FY09.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010
Congressional Add: 1) JMDS	2.373	0.000
<i>FY 2009 Accomplishments:</i> Congressional Interest Item - CATOX system. Developed, conducted testing, and integrated a CATOX system into a U.S. Army vehicle.		
Congressional Adds Subtotals	2.373	0.000

C. Other Program Funding Summary (\$ in Millions)

N/A

D. Acquisition Strategy

JMDS

Joint Material Decontamination System (JMDS) - Congressional Interest Item.

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT CP4: <i>COUNTERPROLIFERATION SUPPORT (ACD&P)</i>

E. Performance Metrics

N/A

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT CP4: <i>COUNTERPROLIFERATION SUPPORT (ACD&P)</i>
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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015							
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4				
** JMDS - Catalytic Oxidation Integrated Demonstration			■	■	■	■	■																									

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT CP4: <i>COUNTERPROLIFERATION SUPPORT (ACD&P)</i>
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Schedule Details

Event	Start		End	
	Quarter	Year	Quarter	Year
** JMDS - Catalytic Oxidation Integrated Demonstration	3	2009	3	2010

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>				R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>				PROJECT DE4: <i>DECONTAMINATION SYSTEMS (ACD&P)</i>			
COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
DE4: <i>DECONTAMINATION SYSTEMS (ACD&P)</i>	4.822	1.792	7.051	0.000	7.051	5.748	1.386	0.000	0.000	Continuing	Continuing
Quantity of RDT&E Articles											

A. Mission Description and Budget Item Justification

This ACD&P project supports the development of decontamination systems utilizing solutions that will remove and/or detoxify contaminated material without damaging combat equipment, personnel, or the environment. Decontamination systems provide a force restoration capability for units that become contaminated. Development efforts will provide systems that reduce operational impact and logistics burden, reduce sustainment costs, increase safety, and minimize environmental effects over currently fielded decontaminants.

This funding supports Human Remains Decontamination System (HRDS) and Joint Platform Interior Decontamination/Joint Service Sensitive Equipment Decontamination (JPID/JSSSED) Congressional Interest Item programs.

The HRDS, Increment I, will utilize mature technologies to provide the capability for safe intra-theater handling and storage of Contaminated Human Remains (CHR) associated with a Chemical Warfare Agent (CWA) event. HRDS will be a Family-of-Systems (FoS) designed to leverage differing technology and requirements readiness across three systems: (1) a Contaminated Human Remains Pouch (CHRP) to support the initial recovery of CHR from Point of Fatality to a Mortuary Affairs Decontamination Collection Point (MADCP); (2) a Contaminated Remains Transfer Case System (CHRTS) capability to store or transport CHR post MADCP operations; and (3) a Remains Decontamination System (RDS) to support the capability to store or transport CHR post MADCP operations. The HRDS will provide the Services the capability to: 1) Safely recover, handle, and transport contaminated human remains prior to decontamination at a MADCP; 2) Enable mortuary affairs units to safely perform their mission with a critical task being that of extensively documenting decedent data and obtaining DNA samples to facilitate positive identification of remains; 3) Fully decontaminate human remains (external), and; 4) Safely allow transport of decontaminated human remains from the MADCP to a final destination in the continental United States for final disposition.

The JPID/JSSSED will fill the capability gap to decontaminate chemical and biological warfare agents from vehicle/aircraft/building interiors, sensitive equipment within, and the associated cargo. This is a new capability that currently does not exist in the DoD. The program goal is to use multiple technologies to provide sensitive equipment and platform interiors decontamination capability.

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT DE4: <i>DECONTAMINATION SYSTEMS (ACD&P)</i>
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The Decontamination Family of Systems (DFoS) program facilitates the rapid transition of mature Science and Technology (S&T) research developments to existing JPM - Decon Program of Record (PoR) and guides S&T community efforts toward meeting the needs of the Warfighter. Leveraging the outcomes of the Analysis of Alternatives scheduled for FY10, DFoS will develop a FoS, to include equipment, to improve decontamination processes and decontaminant solutions to meet the capability gaps for decontaminating NTA and chemical and biological warfare agents from personnel, equipment, vehicle, ship, and aircraft interiors/exterior, terrain and fixed facility interiors/exterior.

Tactical, Cargo, and Rotary Wing Aircraft Decon (Congressional Interest Item): Develop the capability to decontaminate a broad range of military aircraft in the event of a chemical or biological attack.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) DFS <i>FY 2011 Base Plans:</i> Initiate development of non-traditional agent (NTA) efforts to include initial studies and modeling for decontamination assurance spray, chemical decontaminant, reactive skin decontamination lotion/oxime evaluation for NTA decontamination on equipment, effluent decontamination and strippable/sealant coatings.	0.000	0.000	7.051	0.000	7.051
2) HRDS <i>FY 2009 Accomplishments:</i> Developed and refined metrics to support Analysis of Alternatives. Began development of Technology Development Strategy (TDS), Test and Evaluation Strategy (TES) and Systems Engineering Plan (SEP).	1.658	0.000	0.000	0.000	0.000
Accomplishments/Planned Programs Subtotals	1.658	0.000	7.051	0.000	7.051
	FY 2009	FY 2010			
Congressional Add: 1) JMDS	1.582	0.000			

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT DE4: <i>DECONTAMINATION SYSTEMS (ACD&P)</i>
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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010
<i>FY 2009 Accomplishments:</i> Congressional Interest Item - Reactive Overlay and Removable Coatings.		
Congressional Add: 2) JPID <i>FY 2009 Accomplishments:</i> Congressional Interest Item - Environmentally Friendly Aircraft Decontamination System (EFADS). Developed a VHP/Hot Air Prototype Decon System to support the decontamination of Tactical and Cargo Aircraft.	1.582	0.000
Congressional Add: 3) JPID <i>FY 2010 Plans:</i> Congressional Interest Item - Tactical, Cargo, & Rotary Wing Aircraft Decontamination.	0.000	1.792
Congressional Adds Subtotals	3.164	1.792

C. Other Program Funding Summary (\$ in Millions)

Line Item	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	FY 2012	FY 2013	FY 2014	FY 2015	Cost To Complete	Total Cost
• DE5: <i>DECONTAMINATION SYSTEMS (SDD)</i>	16.611	36.786	28.499		28.499	23.944	25.770	14.701	5.928	Continuing	Continuing
• JD0055: <i>JOINT SERVICE PERSONNEL/SKIN DECON SYSTEM (JSPDS)</i>	8.280	4.466	0.000		0.000	0.000	0.000	8.645	9.105	Continuing	Continuing
• JD0056: <i>JS TRANS DECON SYSTEM - SMALL SCALE (JSTDS-SS)</i>	12.124	21.940	18.160		18.160	12.924	7.900	5.455	4.459	Continuing	Continuing
	0.000	0.000	0.000		0.000	4.097	14.064	18.977	25.604	Continuing	Continuing

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT DE4: <i>DECONTAMINATION SYSTEMS (ACD&P)</i>
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C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011 Base</u>	<u>FY 2011 OCO</u>	<u>FY 2011 Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To Complete</u>	<u>Total Cost</u>
• JD0060: <i>JOINT PLATFORM INTERIOR DECON (JPID)</i>											
• JD0061: <i>JS SENSITIVE EQUIP DECON (JSSSED)</i>	0.000	0.000	0.000		0.000	14.648	0.000	0.000	0.000	Continuing	Continuing
• JD0062: <i>HUMAN REMAINS DECON SYSTEM (HRDS)</i>	0.000	0.000	3.410		3.410	3.064	2.936	0.992	0.000	Continuing	Continuing

D. Acquisition Strategy

DFS

DFoS will utilize an incremental acquisition strategy to transition various developmental technology efforts (COTS, JSTO, DTRA efforts, etc.) to fill current and future capability gaps. DFoS will support MDAPs and Programs of Record (POR) by guiding S&T efforts and transitioning mature technologies to meet program requirements. The DFoS acquisition will be managed as a Family-of-Systems (FoS), leveraging differing technologies in each subsystem to fulfill Warfighter capability gaps. A multi-phased Analysis of Alternatives (AoA) will be conducted to identify and evaluate the operational effectiveness of potential material solutions to satisfy Service requirements. As each AoA phase is completed, individual systems and their respective phases of entry will be identified. Industry will be solicited and through competitive prototyping, material solutions will be down-selected for continued development and fielding as a new joint force capability.

HRDS

The Human Remains Decontamination System (HRDS) acquisition will be managed as a Family-of-Systems (FoS), leveraging differing technologies in each subsystem to fulfill Warfighter capability gaps. A multi-phased Analysis of Alternatives (AoA) is being conducted for the HRDS FoS to identify and evaluate the operational effectiveness of potential material solutions to satisfy Service requirements. As each AoA phase is completed, individual systems and their respective phases of entry will be identified. Industry will be solicited and through competitive prototyping, material solutions will be down-selected for continued development and fielding as a new joint force capability.

JMDS

Joint Material Decontamination System (JMDS) - Congressional Interest Item.

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT DE4: <i>DECONTAMINATION SYSTEMS (ACD&P)</i>

E. Performance Metrics

N/A

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT DE4: <i>DECONTAMINATION SYSTEMS (ACD&P)</i>
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Product Development (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** DFS - NTA Decon Assurance Spray	C/FFP	TBD	0.000	0.000		1.000	Jan 2011	0.000		1.000	0.000	1.000	0.000
NTA Chemical Decon	C/FFP	TBD	0.000	0.000		0.850	Jan 2011	0.000		0.850	0.000	0.850	0.000
RSDL/Oxime for NTA Decon on Equipment	C/FFP	TBD	0.000	0.000		1.000	Jan 2011	0.000		1.000	0.000	1.000	0.000
Effluent Decon for NTA Contaminated Run-off	C/FFP	TBD	0.000	0.000		0.859	Jan 2011	0.000		0.859	0.000	0.859	0.000
NTA Strippable / Sealant Coatings	C/FFP	TBD	0.000	0.000		0.950	Jan 2011	0.000		0.950	0.000	0.950	0.000
** JPID - Congressional Interest Item - Tactical, Cargo & Rotary Wing Aircraft Decontamination	SS/FFP	Steris Corp Mentor, Ohio	0.000	1.792	Jan 2010	0.000		0.000		0.000	0.000	1.792	0.000
Subtotal			0.000	1.792		4.659		0.000		4.659	0.000	6.451	0.000

Remarks
DFoS funding increased for NTAs in FY11.

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT DE4: <i>DECONTAMINATION SYSTEMS (ACD&P)</i>
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Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** DFS - IPT Technical Support	MIPR	RDECOM-Natick MA	0.000	0.000		0.400	Jan 2011	0.000		0.400	0.000	0.400	0.000
Subtotal			0.000	0.000		0.400		0.000		0.400	0.000	0.400	0.000

Remarks

Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** DFS - NTA Decon Assurance Spray	MIPR	TBD	0.000	0.000		0.420	Jan 2011	0.000		0.420	0.000	0.420	0.000
NTA Chemical Decon	MIPR	TBD	0.000	0.000		0.320	Jan 2011	0.000		0.320	0.000	0.320	0.000
RSDL/Oxime evaluation for NTA Decon on Equipment	MIPR	TBD	0.000	0.000		0.420	Jan 2011	0.000		0.420	0.000	0.420	0.000
Effluent Decon for NTA Contaminated Run-off	MIPR	TBD	0.000	0.000		0.300	Jan 2011	0.000		0.300	0.000	0.300	0.000
NTA Strippable / Sealant Coatings	MIPR	TBD	0.000	0.000		0.532	Jan 2011	0.000		0.532	0.000	0.532	0.000
Subtotal			0.000	0.000		1.992		0.000		1.992	0.000	1.992	0.000

Remarks

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT DE4: <i>DECONTAMINATION SYSTEMS (ACD&P)</i>
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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
** DFS - NTA Decon Assurance Spray									■	■	■	■	■	■	■	■	■											
NTA Chemical Decon									■	■	■	■	■	■	■	■	■	■	■	■								
RSDL/Oxime evaluation for NTA Decon on Equipment									■	■	■	■	■	■	■	■	■											
Effluent Decon for NTA Contaminated Run-off									■	■	■	■	■	■	■	■	■	■	■	■								
NTA Strippable / Sealant Coatings									■	■	■	■	■	■	■	■	■	■	■	■								
** HRDS - CHRT Market Survey	■																											
HRDS MDD				■																								
HRDS Document Preparation, technical support, and test planning						■	■	■	■	■																		
CHRP/CHRT Development Testing									■	■																		
CHRP/CHRT MS C											■	■																
CHRP/CHRT Fielding												■	■	■	■	■	■	■	■	■	■	■	■	■	■	■		
** JMDS - Catalytic Oxidation Integrated Demonstration			■	■	■	■																						
** JPID - Cong Interest Item - Environmentally Friendly Aircraft Decon System				■	■	■	■																					
Cong Interest Item - Tactical, Cargo & Rotary Wing Aircraft Decontamination						■	■	■	■	■	■	■																

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT DE4: <i>DECONTAMINATION SYSTEMS (ACD&P)</i>
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Schedule Details

Event	Start		End	
	Quarter	Year	Quarter	Year
** DFS - NTA Decon Assurance Spray	2	2011	1	2013
NTA Chemical Decon	2	2011	4	2013
RSDL/Oxime evaluation for NTA Decon on Equipment	2	2011	1	2013
Effluent Decon for NTA Contaminated Run-off	2	2011	4	2013
NTA Strippable / Sealant Coatings	2	2011	4	2013
** HRDS - CHRT Market Survey	1	2009	1	2009
HRDS MDD	4	2009	4	2009
HRDS Document Preparation, technical support, and test planning	2	2010	2	2011
CHRP/CHRT Development Testing	1	2011	2	2011
CHRP/CHRT MS C	3	2011	4	2011
CHRP/CHRT Fielding	4	2011	2	2014
** JMDS - Catalytic Oxidation Integrated Demonstration	3	2009	3	2010
** JPID - Cong Interest Item - Environmentally Friendly Aircraft Decon System	4	2009	4	2010
Cong Interest Item - Tactical, Cargo & Rotary Wing Aircraft Decontamination	2	2010	4	2011

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT IP4: <i>INDIVIDUAL PROTECTION (ACD&P)</i>
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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
IP4: <i>INDIVIDUAL PROTECTION (ACD&P)</i>	0.000	0.000	3.172	0.000	3.172	0.000	0.000	0.000	0.000	Continuing	Continuing
Quantity of RDT&E Articles											

A. Mission Description and Budget Item Justification

This project funds ACD&P of a Lightweight Chemical Biological Ensemble (LCBE) aimed at improving current protection levels while reducing physiological and logistical burdens. The goal is to provide equipment that allows the individual soldier, sailor, airman, or marine to operate in a contaminated chemical and biological (CB) environment with no or minimal degradation to his/her performance.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) LCBE <i>FY 2011 Base Plans:</i> Prepare Request for Information (RFI)/Request for Proposal (RFP); initiate developmental testing (DT) efforts for LCBE Increment 1. Acquire prototypes and perform physical testing and chemical agent testing. Initiates development to reduce thermal burden/bulk/weight over existing CB ensemble, increase cooling/venting potential, improve operational capabilities.	0.000	0.000	3.172	0.000	3.172
Accomplishments/Planned Programs Subtotals	0.000	0.000	3.172	0.000	3.172

C. Other Program Funding Summary (\$ in Millions)

Line Item	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	FY 2012	FY 2013	FY 2014	FY 2015	Cost To Complete	Total Cost
• IP5: <i>INDIVIDUAL PROTECTION (SDD)</i>	18.363	21.094	9.678		9.678	4.833	3.044	0.756	0.563	Continuing	Continuing
	0.000	0.000	0.000		0.000	0.000	0.000	0.000	0.000	Continuing	Continuing

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT IP4: <i>INDIVIDUAL PROTECTION (ACD&P)</i>
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C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011 Base</u>	<u>FY 2011 OCO</u>	<u>FY 2011 Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To Complete</u>	<u>Total Cost</u>
• JSM001: <i>JOINT SERVICE MASK LEAKAGE TESTER (JSMLTS)</i>											
• MA0400: <i>PROTECTIVE CLOTHING (JSLIST)</i>	37.484	20.393	17.887		17.887	18.208	9.429	6.943	6.944	Continuing	Continuing

D. Acquisition Strategy

LCBE

The Lightweight Chemical Biological Ensemble (LCBE) program will pursue an evolutionary incremental approach to provide capability to the Warfighter. Each increment of LCBE will provide technologies with military utility that are modular in function, and offer improvement in form and fit over current systems. The LCBE program will develop, integrate, test, procure and field systems that increase Warfighter operational performance in a CBRN environment via the use of emerging technologies and by leveraging tradespace in areas such as protection level, heat stress, durability, antimicrobial properties, launderability, self-detoxification, protection time, etc. Where appropriate, modeling and simulation tools will be used to lower LCBE program risks, reduce costs and ensure a high confidence in selected technologies.

LCBE INCREMENT 1

The LCBE will use an evolutionary acquisition strategy with phased development. The first LCBE increment will provide an operationally useful and supportable capability in as short a time as possible. Accordingly, Increment 1 of LCBE will incorporate an accelerated development cycle leveraging existing COTS technologies that will, at a minimum, provide a lightweight CB protective garment capability. Gate testing and down-selection of prototypes will comprise the initial phases of the Government's testing program. A competitively awarded contract is planned for DT and Operational Assessment (OA) will occur prior to MS C. Appropriate system requirements reviews, test readiness reviews, producibility reviews and audits will be scheduled as required prior to each milestone.

Future increments of LCBE shall be defined via separate Capability Development Document (CDDs)/Capability Production Document (CPDs) and will follow a similar path/process from MS A or MS B through MS C/FRP and will leverage preceding efforts to the greatest extent possible, maintaining commonality and synergy across all increments.

E. Performance Metrics

N/A

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT IP4: <i>INDIVIDUAL PROTECTION (ACD&P)</i>
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Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** LCBE - ES S - Engineering IPT	MIPR	Various	0.000	0.000		0.600	Jan 2011	0.000		0.600	0.000	0.600	0.000
ES S - Tech Demos	MIPR	Various	0.000	0.000		1.993	Apr 2011	0.000		1.993	0.000	1.993	0.000
Subtotal			0.000	0.000		2.593		0.000		2.593	0.000	2.593	0.000

Remarks

Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** LCBE - PM/MS S - JPM IP Program Management	MIPR	Various	0.000	0.000		0.579	Jan 2011	0.000		0.579	0.000	0.579	0.000
Subtotal			0.000	0.000		0.579		0.000		0.579	0.000	0.579	0.000

Remarks

	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
Project Cost Totals		0.000	0.000		3.172		0.000	3.172	0.000	3.172	0.000

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT IP4: <i>INDIVIDUAL PROTECTION (ACD&P)</i>
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	Total Prior Years Cost	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
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Remarks

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT IP4: <i>INDIVIDUAL PROTECTION (ACD&P)</i>
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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
** LCBE - LCBE Start DT											■	■	■															
LCBE Tech Demo									■	■	■	■	■															

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT IP4: <i>INDIVIDUAL PROTECTION (ACD&P)</i>
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Schedule Details

Event	Start		End	
	Quarter	Year	Quarter	Year
** LCBE - LCBE Start DT	3	2011	1	2012
LCBE Tech Demo	4	2010	1	2012

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT IS4: <i>INFORMATION SYSTEMS (ACD&P)</i>
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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
IS4: <i>INFORMATION SYSTEMS (ACD&P)</i>	0.000	0.000	11.221	0.000	11.221	3.404	4.565	4.676	4.741	Continuing	Continuing
Quantity of RDT&E Articles											

A. Mission Description and Budget Item Justification

This Advanced Component Development and Prototypes (ACD&P) funding supports the Joint Effects Model (JEM) Program.

The JEM is DoD's only accredited model for predicting hazards associated with the release of contaminants into the environment. JEM is being developed in separate increments and is capable of modeling hazards in a variety of scenarios including: counterforce, passive defense, accident and/or incidents (Increment 1); high altitude releases, urban NBC environments (Increment 2); building interiors, and human performance degradation (Increment 3). Battle space commanders and first responders must have a CBRN hazard prediction capability in order to make decisions that will minimize risks of CBRN contamination and enable them to continue mission operations. JEM operates in an integrated fashion with operational and tactical Command, Control, Communications, Computers, Intelligence, Surveillance and Reconnaissance (C4ISR) systems, and in a standalone mode. JEM will interface and communicate with the other programs such as JWARN, JOEF, weather systems, intelligence systems, and various databases.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) JEM Analysis of Alternatives Support <i>FY 2011 Base Plans:</i> Provide Chemical, Biological, Radiological and Nuclear subject matter experts to support the Analysis of Technical Alternatives (ATA).	0.000	0.000	0.689	0.000	0.689
2) JEM	0.000	0.000	0.643	0.000	0.643

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010				
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>		R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>		PROJECT IS4: <i>INFORMATION SYSTEMS (ACD&P)</i>				
B. Accomplishments/Planned Program (\$ in Millions)								
				FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
Analysis of Technical Alternatives (ATA) Evaluation <i>FY 2011 Base Plans:</i> Evaluate and assess results of ATA including a Technology Readiness Assessment of the candidate technologies. Analyze the impact of implementing the emerging technologies in the JEM architecture.								
3) JEM Prototyping <i>FY 2011 Base Plans:</i> Conduct competitive prototyping contracting efforts for JEM Increment 2.				0.000	0.000	4.863	0.000	4.863
4) JEM Increment 2 User Assessments and Demonstrations <i>FY 2011 Base Plans:</i> Prepare for and conduct JEM Increment 2 User Assessments and Demonstrations to validate requirements and system performance, and to evaluate critical science and technology within software prototype(s).				0.000	0.000	1.326	0.000	1.326
5) JEM <i>FY 2011 Base Plans:</i> Conduct government developmental testing and analysis, to include Technology Readiness Assessment of software submitted for evaluation during competitive prototyping.				0.000	0.000	0.961	0.000	0.961
6) JEM Administrative Preparation for Development Contract				0.000	0.000	0.396	0.000	0.396

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT IS4: <i>INFORMATION SYSTEMS (ACD&P)</i>
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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2011 Base Plans:</i> Initiate pre-MS B contractual efforts: develop proposal package, release draft Request for Proposal (RFP), prepare final EMD request for proposal, release RFP, conduct source selection training, conduct source selection and complete proposal evaluations.					
7) JEM Management Support <i>FY 2011 Base Plans:</i> Provide strategic, tactical planning, program/financial management, costing, contracting, scheduling, acquisition oversight, and milestone documentation.	0.000	0.000	1.349	0.000	1.349
8) JEM Technical Support <i>FY 2011 Base Plans:</i> Provide engineering and technical support to Increment 2 development. Provide system independent verification and validation.	0.000	0.000	0.994	0.000	0.994
Accomplishments/Planned Programs Subtotals	0.000	0.000	11.221	0.000	11.221

C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011 Base</u>	<u>FY 2011 OCO</u>	<u>FY 2011 Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To Complete</u>	<u>Total Cost</u>
• G47101: <i>JOINT WARNING & REPORTING NETWORK (JWARN)</i>	4.375	6.551	6.903		6.903	8.078	5.590	8.183	8.423	Continuing	Continuing
	45.694	27.301	13.844		13.844	24.984	24.872	25.345	25.775	Continuing	Continuing

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT IS4: <i>INFORMATION SYSTEMS (ACD&P)</i>
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C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011 Base</u>	<u>FY 2011 OCO</u>	<u>FY 2011 Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To Complete</u>	<u>Total Cost</u>
• IS5: <i>INFORMATION SYSTEMS (SDD)</i>					0.000	0.000	0.000	0.000	0.000	Continuing	Continuing
• IS6: <i>INFORMATION SYSTEMS (RDT&E MGT SUPPORT)</i>	0.000	0.000	0.000		0.000	0.000	0.000	0.000	0.000	Continuing	Continuing
• IS7: <i>INFORMATION SYSTEMS (OP SYS DEV)</i>	0.897	1.302	1.821		1.821	1.669	1.518	1.555	1.577	Continuing	Continuing
• JC0208: <i>JOINT EFFECTS MODEL (JEM)</i>	5.546	3.482	3.482		3.482	0.000	0.000	3.369	3.568	Continuing	Continuing
• JC0209: <i>JOINT OPERATIONAL EFFECTS FEDERATION (JOEF)</i>	0.000	0.000	0.000		0.000	0.000	2.482	2.480	3.716	Continuing	Continuing

D. Acquisition Strategy

JEM

The Joint Effects Model (JEM) is following an evolutionary acquisition approach that will allow rapid fielding of existing technologies while further research and development (R&D) continues in order to mature the technologies required for subsequent versions of JEM. It is now being fielded in increments of capabilities. Each increment will retain the functionality of the preceding increment. The JEM development effort will be aligned with the evolving JPEO-CBD architectures and technologies, as well as with Service C2 systems. JEM is expected to develop three distinct increments of software. JEM will define and publish its web-services interface; the JEM interface will be the same on all systems, utilizing data definitions from the approved CBRN data model as appropriate. A cost plus award fee contract was awarded for the follow-on JEM contract for integration and development.

E. Performance Metrics

N/A

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT IS4: <i>INFORMATION SYSTEMS (ACD&P)</i>
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Product Development (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** JEM - SW SB - JEM	MIPR	San Diego CA	0.000	0.000		7.521	Jan 2011	0.000		7.521	0.000	7.521	0.000
Subtotal			0.000	0.000		7.521		0.000		7.521	0.000	7.521	0.000

Remarks

Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** JEM - ES S - JEM System	MIPR	Various	0.000	0.000		0.994	Jan 2011	0.000		0.994	0.000	0.994	0.000
Subtotal			0.000	0.000		0.994		0.000		0.994	0.000	0.994	0.000

Remarks

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT IS4: <i>INFORMATION SYSTEMS (ACD&P)</i>
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Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** JEM - DTE S - JEM	MIPR	Various	0.000	0.000		0.961	Jan 2011	0.000		0.961	0.000	0.961	0.000
Subtotal			0.000	0.000		0.961		0.000		0.961	0.000	0.961	0.000

Remarks

Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** JEM - PM/MS S - JEM	MIPR	Various	0.000	0.000		1.745	Jan 2011	0.000		1.745	0.000	1.745	0.000
Subtotal			0.000	0.000		1.745		0.000		1.745	0.000	1.745	0.000

Remarks

Project Cost Totals	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
		Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
	0.000	0.000		11.221		0.000		11.221	0.000	11.221	0.000

Remarks

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT IS4: <i>INFORMATION SYSTEMS (ACD&P)</i>
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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
** JEM - Increment 1 - Pre-planned Product Improvement (P3I)	■	■	■	■	■	■	■	■	■	■	■																	
Increment 1 - Production and Deployment	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■												
Increment 1 - Developmental Maintenance	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■												
Increment 1 - Follow-on Test and Evaluation				■	■	■																						
Increment 2 - Material Development Decision (MDD)					■																							
Increment 2 - Technology Development	■	■	■	■	■	■	■	■	■	■																		
Increment 2 - Analysis of Alternatives					■	■	■	■	■																			
Increment 2 - DT (Cont)							■	■	■	■	■	■	■	■	■	■	■	■										
Increment 2 - DT (Gov't)							■	■	■	■	■	■	■	■	■	■	■	■	■	■								
Increment 2 - User Assessments									■	■	■	■																
Increment 2 - Milestone A (MS A)									■																			
Increment 2 - Prototype Development/Testing									■	■	■	■																
Increment 2 - Capability Development Document (CDD)									■	■	■																	
Increment 2 - Engineering and Manufacturing Development									■	■	■	■	■	■														
Increment 2 - Milestone B (MS B)													■															
Increment 2 - Capability Production Document (CPD)													■	■	■													
Increment 2 - Operational Assessment (OA)																■												

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT IS4: <i>INFORMATION SYSTEMS (ACD&P)</i>
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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
Increment 2 - Milestone C (MS C)																												
Increment 2 - Multi-Service Operational Test and Evaluation (MOT&E)/LOG Demo																												
Increment 2 - Standalone Full Rate Production (FRP)																												

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT IS4: <i>INFORMATION SYSTEMS (ACD&P)</i>
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Schedule Details

Event	Start		End	
	Quarter	Year	Quarter	Year
** JEM - Increment 1 - Pre-planned Product Improvement (P3I)	3	2008	3	2011
Increment 1 - Production and Deployment	4	2007	4	2012
Increment 1 - Developmental Maintenance	3	2008	4	2012
Increment 1 - Follow-on Test and Evaluation	4	2009	2	2010
Increment 2 - Material Development Decision (MDD)	1	2010	1	2010
Increment 2 - Technology Development	3	2008	2	2011
Increment 2 - Analysis of Alternatives	1	2010	1	2011
Increment 2 - DT (Cont)	3	2010	2	2013
Increment 2 - DT (Gov't)	3	2010	3	2013
Increment 2 - User Assessments	4	2010	3	2011
Increment 2 - Milestone A (MS A)	1	2011	1	2011
Increment 2 - Prototype Development/Testing	1	2011	4	2011
Increment 2 - Capability Development Document (CDD)	1	2011	3	2011
Increment 2 - Engineering and Manufacturing Development	1	2011	2	2012
Increment 2 - Milestone B (MS B)	2	2012	2	2012
Increment 2 - Capability Production Document (CPD)	2	2012	4	2012
Increment 2 - Operational Assessment (OA)	4	2012	4	2012
Increment 2 - Milestone C (MS C)	2	2013	2	2013

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT IS4: <i>INFORMATION SYSTEMS (ACD&P)</i>
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Event	Start		End	
	Quarter	Year	Quarter	Year
Increment 2 - Multi-Service Operational Test and Evaluation (MOT&E)/LOG Demo	4	2013	4	2013
Increment 2 - Standalone Full Rate Production (FRP)	2	2014	2	2014

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>				R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>				PROJECT MB4: <i>MEDICAL BIOLOGICAL DEFENSE (ACD&P)</i>			
COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
MB4: <i>MEDICAL BIOLOGICAL DEFENSE (ACD&P)</i>	7.910	102.437	136.975	0.000	136.975	130.718	131.347	115.985	113.566	Continuing	Continuing
Quantity of RDT&E Articles											

A. Mission Description and Budget Item Justification

The Joint Vaccine Acquisition Program (JVAP) under Chemical Biological Medical Systems (CBMS) funds the technology development phase for vaccines that are directed against validated biological warfare (BW) weapons to include bacteria, viruses, and toxins of biological origin. Effective medical countermeasures to negate the threat of these BW agents are urgently needed. Vaccines have been identified as the most efficient countermeasure against the validated threat of BW weapons. Efforts for medical biological defense product development involve manufacturing process development, formulation development, non-clinical studies, pilot lot manufacturing, Investigational New Drug (IND) application submission and Phase 1 clinical human safety studies. Vaccine effectiveness, will be evaluated in animals to satisfy the requirements of the FDA's "Animal Rule". Products under development in this budget item include Filovirus vaccine.

The Transformational Medical Technologies Initiative (TMTI) was launched to respond to the threat of emerging or intentionally bioengineered biological threats. TMTI's mission is to protect the Warfighter from genetically engineered biological threats by providing a rapid response capability from identification of pathogens to the delivery of medical countermeasures. This mission is accomplished by developing broad spectrum (multi-agent) therapeutics against biological warfare (BW) agents (e.g. one drug that treats multiple agents). The development of broad spectrum therapeutics involves developing a capability to treat exposure to hemorrhagic fever viruses (HFVs) (e.g. Ebola virus) and intracellular bacterial pathogens (ICBs) (e.g. Tularemia). Efforts are further classified as host-directed therapeutics (e.g. drugs that target common pathways within a human to prevent or treat a variety of diseases) or pathogen-directed therapeutics (e.g. drugs that attack a common pathway found in multiple threat agents). Attrition is high throughout the drug development process. Less than 10% of all preclinical compounds become a licensed drug. Causes for attrition include scientific failures, Food and Drug Administration (FDA) rejection at major milestone reviews, and loss through down-selection at DoD Milestone Decision points. The development of medical countermeasures is an arduous process that requires extensive interaction with the FDA, from pre-clinical research to safety tests in human subjects (Phase 1 clinical studies), efficacy tests in humans/animals (Phase 2 clinical studies or pivotal animal efficacy studies), and expanded safety or efficacy studies (Phase 3 clinical studies), which culminate with a request to the FDA to license, market, and produce a drug. This interaction between the Department of Defense (DoD) and the FDA results in a coordinated, unified, and safe effort.

B. Accomplishments/Planned Program (\$ in Millions)

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program			DATE: February 2010			
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT MB4: <i>MEDICAL BIOLOGICAL DEFENSE (ACD&P)</i>				
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2010 Plans:</i> Prepare supporting acquisition documentation, conduct Milestone A, and enter into Advanced Component Development and Prototypes.						
4) JVAP - Filovirus Vaccine <i>FY 2010 Plans:</i> Prepare solicitation documentation and conduct pre-solicitation conference. <i>FY 2011 Base Plans:</i> Release solicitation documentation and conduct source selection to award contract.		0.000	0.200	0.100	0.000	0.100
5) JVAP - Filovirus Vaccine <i>FY 2010 Plans:</i> Initiate non-clinical studies through Interagency Agreements. Initiate procedures for safeguarding biological select agents and toxins. <i>FY 2011 Base Plans:</i> Continue non-clinical studies through Interagency Agreements. Continue procedures for safeguarding biological select agents and toxins.		0.000	10.569	9.108	0.000	9.108
6) JVAP - Filovirus Vaccine <i>FY 2011 Base Plans:</i> Initiate small-scale manufacturing process development.		0.000	0.000	7.410	0.000	7.410
7) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.		0.000	1.264	0.000	0.000	0.000

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program			DATE: February 2010			
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT MB4: <i>MEDICAL BIOLOGICAL DEFENSE (ACD&P)</i>				
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
Accomplishments/Planned Programs Subtotals		0.000	100.844	136.975	0.000	136.975
		FY 2009	FY 2010			
Congressional Add: 1) Vacuum Sampling Pathogen Collection and Concentration <i>FY 2009 Accomplishments:</i> Congressional Interest Item - Vacuum Sampling Pathogen Collection and Concentration. Continued development of the M-Vac System, a field pathogen collection system, along with the Bacteria Reduction System (BRS) to enhance the current capability to find, extract and elute potentially deadly pathogens from unique surfaces. A tube sampler was developed, which enable customers and inspectors to reach and sample inaccessible surfaces and areas. An effort began to miniaturize the sampling platform in order to meet DoD requirements.		3.164	0.000			
Congressional Add: 2) Broad Spectrum Therapeutic Countermeasure <i>FY 2010 Plans:</i> Congressional Interest Item - Broad Spectrum Therapeutic Countermeasure to Organophosphorus Nerve Agents. Initiate development of a broad-spectrum therapeutic capable of protecting both the central and peripheral nervous systems from injury by nerve agents and reducing reliance on pretreatments.		0.000	1.593			
Congressional Add: 3) CRP <i>FY 2009 Accomplishments:</i> Congressional Interest Item - Biological Threat Antibody Research.		1.582	0.000			
		0.791	0.000			

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT MB4: <i>MEDICAL BIOLOGICAL DEFENSE (ACD&P)</i>
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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010
Congressional Add: 4) TT Bio <i>FY 2009 Accomplishments:</i> Congressional Interest Item - IM Formulation Development of Anthrax Therapeutic.		
Congressional Add: 5) TT Bio <i>FY 2009 Accomplishments:</i> Congressional Interest Item - Recombinant Bche Formulation Program.	1.582	0.000
Congressional Add: 6) TT Bio <i>FY 2009 Accomplishments:</i> Congressional Interest Item - Large scale single-use bio reactor for rapid response to bioterrorism.	0.791	0.000
Congressional Adds Subtotals	7.910	1.593

C. Other Program Funding Summary (\$ in Millions)

Line Item	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	FY 2012	FY 2013	FY 2014	FY 2015	Cost To Complete	Total Cost
• JM0001: <i>JOINT BIO AGENT IDENT AND DIAG SYSTEM (JBAIDS)</i>	0.479	0.000	5.571		5.571	0.000	0.000	0.000	0.000	Continuing	Continuing
• JX0005: <i>DOD BIOLOGICAL VACCINE PROCUREMENT</i>	38.109	12.701	12.824		12.824	3.385	3.466	56.416	98.759	Continuing	Continuing
• JX0210: <i>CRITICAL REAGENTS PROGRAM (CRP)</i>	0.000	0.000	0.994		0.994	0.993	0.993	0.992	0.000	Continuing	Continuing
• MB5: <i>MEDICAL BIOLOGICAL DEFENSE (SDD)</i>	87.676	57.558	141.680		141.680	161.732	159.144	141.481	111.671	Continuing	Continuing
	180.425	203.723	115.233		115.233	125.666	109.737	115.049	117.289	Continuing	Continuing

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT MB4: <i>MEDICAL BIOLOGICAL DEFENSE (ACD&P)</i>
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C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011</u> <u>Base</u>	<u>FY 2011</u> <u>OCO</u>	<u>FY 2011</u> <u>Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To</u> <u>Complete</u>	<u>Total Cost</u>
• TB3: <i>MEDICAL BIOLOGICAL DEFENSE (ATD)</i>											

D. Acquisition Strategy

CRP

The Critical Reagents Program's (CRP) strategy establishes a core research and development capability to develop biological threat agent, genomic reference materials (antigens, nucleic acids, and antibodies) and detection and diagnostic assays for biothreat agent detection that shall be horizontally inserted across multiple detection and diagnostic platforms. In addition, this strategy will implement a formal, validated advanced development process to transition new assays into production and integration with the appropriate detection/diagnostic platform.

TMTI

Transformational Medical Technology Initiative's (TMTI) ultimate goal is the delivery of FDA-licensed, therapeutics to the Warfighter. This goal can be reached through any one of the following three acquisition approaches: 1) through the discovery of new drugs; 2) through application of new drug indications (i.e., through a commercial off-the-shelf (COTS) approach); or, 3) through the re-engineering of previously developed drugs (i.e., through a Modified COTS approach). This may involve FDA-approved drugs or previously developed drug compounds that do not have an FDA license. Each of these approaches will require different entry points into both the drug development process and the defense acquisition management timeline. Moreover, each of these approaches will likely experience a different set of FDA regulatory requirements. In order to execute the overall acquisition strategy, TMTI has partnered with other elements within the DoD Chemical and Biological Defense Program, DoD agencies, private industry, and other DoD laboratories for the development of TMTI products. The contract types used to execute the program will depend on the circumstances, including maturity of the science, the legalities surrounding Intellectual Property (IP) and patent rights, and even the size of the performer. Cost Plus Fixed Fee and Cost Plus Incentive Fee contracts will be used with traditional or nontraditional defense contractors for most advanced development contracts. Finally, developing platform technologies, such as modeling and simulation to predict drug-to-drug interaction effects prior to actual clinical trials, and the use of genetic sequencing and a bioinformatics backbone, are examples of how TMTI managers intend to augment private industry best practices to streamline the program management, test and evaluation, and overall TMTI product development.

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program	DATE: February 2010
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APPROPRIATION/BUDGET ACTIVITY	R-1 ITEM NOMENCLATURE	PROJECT
0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	MB4: <i>MEDICAL BIOLOGICAL DEFENSE (ACD&P)</i>

Contract will be competed at Milestone A. Developer(s) and candidate(s) will be selected based on existing test data, technical approach and business case considerations. Because of inherent risk in development of vaccines for viral hemorrhagic fevers, funding requested will support development of two vaccine candidates through Milestone B, at which time, a down-select to a single candidate will be conducted. If technologically feasible, a combined Ebola/Marburg vaccine will be developed. In addition to efforts conducted under a contract, portions of the non-clinical work will be conducted under an Interagency Agreement with the US Army Medical Research Institute of Infectious Diseases (USAMRIID), because of the limited number of Biological Safety Level (BSL) 4 facilities.

E. Performance Metrics

N/A

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT MB4: <i>MEDICAL BIOLOGICAL DEFENSE (ACD&P)</i>
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Product Development (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** TMTI - SW SB - Therapeutic validation Contract #1	C/CPIF	TBD	0.000	5.257	Apr 2010	5.765	Apr 2011	0.000		5.765	0.000	11.022	0.000
SW SB - Therapeutic validation Contract #2	C/CPIF	TBD	0.000	5.256	Apr 2010	5.764	Apr 2011	0.000		5.764	0.000	11.020	0.000
SW SB - Therapeutic validation Contract #3	C/CPIF	TBD	0.000	5.257	Apr 2010	5.764	Apr 2011	0.000		5.764	0.000	11.021	0.000
SW SB - Therapeutic validation Contract #4	C/CPIF	TBD	0.000	5.256	Apr 2010	5.765	Apr 2011	0.000		5.765	0.000	11.021	0.000
SW SB - Therapeutic validation Contract #5	C/CPIF	TBD	0.000	5.256	Apr 2010	5.764	Apr 2011	0.000		5.764	0.000	11.020	0.000
SW SB - Therapeutic validation Contract #6	C/CPIF	TBD	0.000	5.256	Apr 2010	5.765	Apr 2011	0.000		5.765	0.000	11.021	0.000
SW S - Technology Contract	C/CPIF	TBD	0.000	0.000		7.635	Oct 2010	0.000		7.635	0.000	7.635	0.000
** VAC FILO - Manufacturing, Validation, Pilot Lot, and Consistency Lot Production	C/CPIF	TBD	0.000	0.000		4.398	Jan 2011	0.000		4.398	0.000	4.398	0.000
Non Clinical Studies	MIPR	USAMRIID Fort Detrick, MD	0.000	10.569	Apr 2010	3.250	Jan 2011	0.000		3.250	0.000	13.819	0.000
Subtotal			0.000	42.107		49.870		0.000		49.870	0.000	91.977	0.000

Remarks

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT MB4: <i>MEDICAL BIOLOGICAL DEFENSE (ACD&P)</i>
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Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** TMTI - TD/D SB - Regulatory Integration, Quality Assurance, & FDA Support Efforts - Contract #1	C/CPIF	TBD	0.000	3.754	Apr 2010	4.117	Apr 2011	0.000		4.117	0.000	7.871	0.000
TD/D SB - Regulatory Integration, Quality Assurance, & FDA Support Efforts - Contract #2	C/CPIF	TBD	0.000	3.755	Apr 2010	4.118	Apr 2011	0.000		4.118	0.000	7.873	0.000
ES C - Regulatory Integration, Quality Assurance, & FDA Support Efforts - Contract #3	C/CPIF	TBD	0.000	3.754	Apr 2010	4.117	Apr 2011	0.000		4.117	0.000	7.871	0.000
ES C - Regulatory Integration, Quality Assurance, & FDA Support Efforts - Contract #4	C/CPIF	TBD	0.000	3.755	Apr 2010	4.118	Apr 2011	0.000		4.118	0.000	7.873	0.000
ES C - Regulatory Integration, Quality Assurance, & FDA Support Efforts - Contract #5	C/CPIF	TBD	0.000	3.755	Apr 2010	4.117	Apr 2011	0.000		4.117	0.000	7.872	0.000
ES C - Regulatory Integration, Quality Assurance, & FDA	C/CPIF	TBD	0.000	3.754	Apr 2010	4.117	Apr 2011	0.000		4.117	0.000	7.871	0.000

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APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT MB4: <i>MEDICAL BIOLOGICAL DEFENSE (ACD&P)</i>
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Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
Support Efforts - Contract #6													
TD/D C - Technologies	C/CPIF	TBD	0.000	0.000		5.453	Oct 2010	0.000		5.453	0.000	5.453	0.000
** VAC FILO - Regulatory Integration (Environmental and FDA Documentation) and Delivery System	C/CPIF	TBD	0.000	0.000		2.288	Jan 2011	0.000		2.288	0.000	2.288	0.000
Subtotal			0.000	22.527		32.445		0.000		32.445	0.000	54.972	0.000

Remarks

Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** CONG - Congressional Interest Item - Broad Spectrum Therapeutic	SS/FFP	TBD	0.000	1.513	Apr 2010	0.000		0.000		0.000	0.000	1.513	0.000
** TMTI - DTE C - Phase I trials - Contract #1	C/CPIF	TBD	0.000	6.007	Apr 2010	4.588	Apr 2011	0.000		4.588	0.000	10.595	0.000
	C/CPIF	TBD	0.000	6.007	Apr 2010	4.588	Apr 2011	0.000		4.588	0.000	10.595	0.000

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APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT MB4: <i>MEDICAL BIOLOGICAL DEFENSE (ACD&P)</i>
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Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
DTE C - Phase I trials - Contract #2													
DTE C - Phase I trials - Contract #3	C/CPIF	TBD	0.000	6.007	Apr 2010	4.588	Apr 2011	0.000		4.588	0.000	10.595	0.000
DTE C - Phase I trials - Contract #4	C/CPIF	TBD	0.000	6.007	Apr 2010	4.588	Apr 2011	0.000		4.588	0.000	10.595	0.000
DTE C - Phase I trials - Contract #5	C/CPIF	TBD	0.000	6.009	Apr 2010	4.588	Apr 2011	0.000		4.588	0.000	10.597	0.000
DTE C - Phase I trials - Contract #6	C/CPIF	TBD	0.000	4.509	Apr 2010	4.588	Apr 2011	0.000		4.588	0.000	9.097	0.000
OTHT C - Technologies	C/CPIF	TBD	0.000	0.000		8.450	Apr 2011	0.000		8.450	0.000	8.450	0.000
DTE C - Phase I trials - IBP Contract #1	C/CPIF	TBD	0.000	0.000		3.000	Apr 2011	0.000		3.000	0.000	3.000	0.000
DTE C - Phase I trials - IBP Contract #2	C/CPIF	TBD	0.000	0.000		3.000	Apr 2011	0.000		3.000	0.000	3.000	0.000
DTE C - Phase I trials - IBP Contract #3	C/CPIF	TBD	0.000	0.000		3.000	Apr 2011	0.000		3.000	0.000	3.000	0.000
DTE C - Phase I trials - IBP Contract #4	C/CPIF	TBD	0.000	0.000		3.000	Apr 2011	0.000		3.000	0.000	3.000	0.000
** VAC FILO - Testing, Evaluation, and Clinical Trials	C/CPIF	TBD	0.000	0.000		4.958	Jan 2011	0.000		4.958	0.000	4.958	0.000
Subtotal			0.000	36.059		52.936		0.000		52.936	0.000	88.995	0.000

Remarks

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT MB4: <i>MEDICAL BIOLOGICAL DEFENSE (ACD&P)</i>
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Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** CONG - Congressional Interest Item - Broad Spectrum Therapeutic	Allot	TBD	0.000	0.080	Apr 2010	0.000		0.000		0.000	0.000	0.080	0.000
** VAC FILO - PM/MS S - Program Management/ Program Manager Support	Allot	CBMS Frederick, MD	0.000	0.200	Jul 2010	1.035	Jul 2011	0.000		1.035	0.000	1.235	0.000
PM/MS S - Contractor Systems Engineering/ Program Management Support	SS/FFP	Goldbelt Raven LLC, Frederick	0.000	0.200	Apr 2010	0.689	Apr 2011	0.000		0.689	0.000	0.889	0.000
** ZSBIR - SBIR/STTR - Aggregated from ZSBIR-SBIR/STTR	MIPR	HQ AMC, Alexandria	0.000	1.264		0.000		0.000		0.000	0.000	1.264	0.000
Subtotal			0.000	1.744		1.724		0.000		1.724	0.000	3.468	0.000

Remarks

	Total Prior Years Cost	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
Project Cost Totals	0.000	102.437	136.975	0.000	136.975	0.000	239.412	0.000

Remarks

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT MB4: <i>MEDICAL BIOLOGICAL DEFENSE (ACD&P)</i>
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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
** CONG - Tactical, Cargo & Rotary Wing Aircraft Decon							■	■	■	■	■	■	■	■	■	■												
** TMTI - Milestone A Decision (Hemorrhagic Fever Virus MCM)			■																									
Contract #1-6 (HFV) Phase I trials							■	■	■	■	■	■																
Milestone A Decision (Intracellular Bacteria Pathogen MCM)								■																				
Contract 1-4 (IBP) Phase I Trials											■	■	■	■	■	■												
Milestone B Decision (Hemorrhagic Fever Viruses)											■																	
** TT Bio - Large Scale Single Use Bio Reactor for Rapid Response to Bioterrorism			■	■																								
Biological Threat Antibody Research			■	■																								
Recombinant Bche Formulation Program			■	■																								
IM Formulation development of Anthrax Therapeutic			■	■																								
** VAC FILO - Prepare Acquisition Documentation						■	■	■	■																			
Prepare and release solicitation						■	■	■	■	■																		
VAC FILO - Milestone A								■																				
VAC FILO - Manufacturing Process Development - Small Scale											■	■	■	■	■	■	■	■	■	■	■							

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT MB4: <i>MEDICAL BIOLOGICAL DEFENSE (ACD&P)</i>
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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
VAC FILO - Non-clinical Studies							■	■	■	■	■	■	■	■	■	■	■	■	■	■	■							

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT MB4: <i>MEDICAL BIOLOGICAL DEFENSE (ACD&P)</i>
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Schedule Details

Event	Start		End	
	Quarter	Year	Quarter	Year
** CONG - Tactical, Cargo & Rotary Wing Aircraft Decon	3	2010	4	2012
** TMTI - Milestone A Decision (Hemorrhagic Fever Virus MCM)	4	2009	4	2009
Contract #1-6 (HFV) Phase I trials	3	2010	4	2011
Milestone A Decision (Intracellular Bacteria Pathogen MCM)	4	2010	4	2010
Contract 1-4 (IBP) Phase I Trials	3	2011	4	2012
Milestone B Decision (Hemorrhagic Fever Viruses)	3	2011	3	2011
** TT Bio - Large Scale Single Use Bio Reactor for Rapid Response to Bioterrorism	3	2009	4	2009
Biological Threat Antibody Research	3	2009	4	2009
Recombinant Bche Formulation Program	3	2009	4	2009
IM Formulation development of Anthrax Therapeutic	3	2009	4	2009
** VAC FILO - Prepare Acquisition Documentation	1	2010	4	2010
Prepare and release solicitation	1	2010	2	2011
VAC FILO - Milestone A	4	2010	4	2010
VAC FILO - Manufacturing Process Development - Small Scale	3	2011	1	2014
VAC FILO - Non-clinical Studies	3	2010	1	2014

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program								DATE: February 2010			
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>				R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>				PROJECT MC4: <i>MEDICAL CHEMICAL DEFENSE (ACD&P)</i>			
COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
MC4: <i>MEDICAL CHEMICAL DEFENSE (ACD&P)</i>	19.365	9.438	0.000	0.000	0.000	2.973	3.661	5.035	14.670	Continuing	Continuing
Quantity of RDT&E Articles											

A. Mission Description and Budget Item Justification

This project funds Advanced Component Development and Prototypes (ACD&P) of countermeasures for chemical agents including life support equipment, diagnostic equipment, prophylactic and therapeutic drugs, and individual/casualty decontamination compounds. A system of medical defense against chemical agents is required to provide protection, to sustain performance in a chemical environment, and to provide for self-aid/buddy-aid and medical treatment of chemical casualties. Fielding of prophylactic and therapeutic drugs requires Food and Drug Administration (FDA) approval. Multiple long-term studies are required to obtain FDA approval resulting in longer program timelines and greater program cost than other non-pharmaceutical product programs. Efficacy testing of most candidate drugs against chemical warfare (CW) agents cannot be conducted in humans; therefore, animal surrogate models must be developed. The program currently funds the: (1) Bioscavenger Increment 2 (BSCAV Inc. 2), which will be used as a prophylaxis against nerve agents; (2) Inhalational Atropine (IA), which will be used to treat continuing nerve agent induced effects after the patient has been evacuated to a medical treatment facility; and (3) Improved Nerve Agent Treatment System (INATS), which will be used as a treatment for nerve agent intoxication to include new indications for Pyridostigmine Bromide (PB) that will be integrated with current therapeutic regimens. Time Temperature Indicators (TTI), Item Unique Identification (IUID), and Radio-Frequency Identification (RFID) will be part of the development effort for incorporation on all medical countermeasures being developed by CBMS-MITS. A TTI is a human readable tab that will provide the Warfighter immediate knowledge if the product is still useable or not. IUID and RFID labels placed on the product will improve inventory management and strategic purchasing, and enable reliable visibility, and capability-based operational readiness.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) BSCAV Increment 2 <i>FY 2009 Accomplishments:</i> Completed Phase 1 clinical safety studies and prepared documentation for Milestone B in 1QFY10.	4.686	0.000	0.000	0.000	0.000
2) BSCAV Increment 2	4.050	0.000	0.000	0.000	0.000

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT MC4: <i>MEDICAL CHEMICAL DEFENSE (ACD&P)</i>				
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2009 Accomplishments:</i> Continued large scale manufacturing, process development, and assay validation. Prepared for transition to Engineering and Manufacturing Development phase.						
3) BSCAV Increment 2 <i>FY 2009 Accomplishments:</i> Completed NTA studies at US Army Medical Research Institute of Chemical Defense (USAMRICD).		0.400	0.000	0.000	0.000	0.000
4) Inhalational Atropine <i>FY 2010 Plans:</i> Initiate process development and cGMP requirements.		0.000	0.800	0.000	0.000	0.000
5) Inhalational Atropine <i>FY 2010 Plans:</i> Initiate formulation, analytical methods and device development and transition to Engineering Manufacturing and Development phase.		0.000	1.167	0.000	0.000	0.000
6) INATS <i>FY 2009 Accomplishments:</i> Completed Investigational New Drug (IND) application effort. <i>FY 2010 Plans:</i> Initiate and complete IND amendment.		0.630	0.552	0.000	0.000	0.000
7) INATS		3.750	3.411	0.000	0.000	0.000

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2009 Accomplishments:</i> Continued process development and small scale Current Good Manufacturing Practices (cGMP) efforts.</p> <p><i>FY 2010 Plans:</i> Complete process development and small scale cGMP efforts.</p>					
<p>8) INATS</p> <p><i>FY 2009 Accomplishments:</i> Continued efficacy, safety, and toxicology studies of candidate oximes.</p> <p><i>FY 2010 Plans:</i> Complete efficacy, safety, and toxicology studies of candidate oximes.</p>	5.849	3.390	0.000	0.000	0.000
<p>9) SBIR</p> <p><i>FY 2010 Plans:</i> Small Business Innovative Research.</p>	0.000	0.118	0.000	0.000	0.000
Accomplishments/Planned Programs Subtotals					
	19.365	9.438	0.000	0.000	0.000

C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011 Base</u>	<u>FY 2011 OCO</u>	<u>FY 2011 Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To Complete</u>	<u>Total Cost</u>
• JM6500: <i>INHALATIONAL ATROPINE (IA)</i>	0.000	0.000	0.000		0.000	0.000	0.000	0.496	0.991	Continuing	Continuing
• JM6555: <i>IMPROVED NERVE AGENT TREATMENT SYSTEM (INATS)</i>	0.000	0.000	0.000		0.000	0.000	0.000	3.966	4.954	Continuing	Continuing

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT MC4: <i>MEDICAL CHEMICAL DEFENSE (ACD&P)</i>
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C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011</u> <u>Base</u>	<u>FY 2011</u> <u>OCO</u>	<u>FY 2011</u> <u>Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To</u> <u>Complete</u>	<u>Total Cost</u>
• JM6677: <i>ADVANCED ANTICONVULSANT SYSTEM (AAS)</i>	0.000	0.000	0.000		0.000	0.000	4.466	9.126	5.187	Continuing	Continuing
• MC5: <i>MEDICAL CHEMICAL DEFENSE (SDD)</i>	14.203	14.027	51.856		51.856	47.835	28.771	12.122	8.171	Continuing	Continuing

D. Acquisition Strategy

BSCAV

The Bioscavenger acquisition strategy consists of a developmental program with three distinct increments.

Increment 1 is butyrylcholinesterase purified from human plasma (i.e., plasma-derived Bioscavenger or pBioscavenger). The Medical Identification and Treatment Systems (MITS) Joint Product Management Office exercises management oversight, and a commercial partner serves as the system integrator during the Technology Development Phase, which includes small scale manufacturing, pre-clinical animal studies, Investigational New Drug (IND) application, and Phase 1 human clinical safety studies.

The Bioscavenger Increment 2 strategy includes a proof-of-concept study followed by an initial down-selection between two different technologies: Recombinant human butyrylcholinesterase (rHuBChE) and small synthetic molecule, awarded to two different contractors. The chosen technology, rHuBChE, will continue to a formal down-selection with the plasma-derived Bioscavenger at Milestone B prior to transition to the Engineering and Manufacturing Development (EMD) phase. Following Milestone B into EMD, MITS will continue to exercise management oversight with system integration support of a commercial partner to ensure manufacturing of the product is in accordance with Food and Drug Administration (FDA) regulations and guidelines. Prior to FDA licensure, the commercial partner will perform a Phase 2 human clinical safety study, definitive animal efficacy studies, and toxicology studies. The SDD phase will culminate in obtaining FDA licensure of the Bioscavenger. During the Production and Deployment phase, the MITS JPMO, in conjunction with a commercial partner, will pursue full rate and stockpile production and conduct any FDA-mandated post-marketing surveillance.

Unlike Bioscavenger Increment 1 and 2 technology, where the Bioscavenger becomes ineffective after binding with nerve agents, Increment 3 will include products that continuously degrade nerve agents while retaining their effectiveness (catalytic Bioscavenger). Because the technology for Increment 3 is immature, a candidate product will not be ready for transition to advanced development until late in the next decade. Therefore, CBMS MITS is exploring alternative technologies to reduce the costs of producing Bioscavenger Increment 2.

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
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<p>IA</p> <p>Medical Identification and Treatment Systems (MITS) Joint Product Management Office (JPMO) will manage the development of Inhalational Atropine (IA) for the DoD. For this Advanced Development effort, the competitively selected contractor will serve as the systems integrator throughout development and shall be responsible for conducting the activities associated with drug development in a manner consistent with eventual approval by the Food and Drug Administration (FDA), including: human clinical safety studies; pharmacokinetic studies; and validated manufacturing. The contractor shall sponsor the drug product to the FDA and hold all approvals and/or licenses.</p> <p>INATS</p> <p>Medical Identification and Treatment Systems (MITS) Joint Product Management Office will serve as the system integrator during the Technology Development Phase that includes pre-clinical animal studies and Phase 1 human clinical safety studies. After Milestone B, during the System Development and Demonstration Phase, MITS and/or a commercial partner (product dependent) will serve as the system integrator to ensure that products are manufactured in accordance with Food and Drug Administration (FDA) regulations and guidelines, appropriate Phase 2 human clinical safety and definitive animal efficacy studies are conducted, and required toxicology studies are performed. During the Production and Deployment Phase, FDA approval will be obtained and full rate and stockpile production will be pursued. Any FDA mandated post-marketing surveillance will be conducted.</p> <p><u>E. Performance Metrics</u> N/A</p>		

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT MC4: <i>MEDICAL CHEMICAL DEFENSE (ACD&P)</i>
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Product Development (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** IA - cGMP Manufacturing requirements	C/CPIF	TBD	0.000	0.767	Jan 2010	0.000		0.000		0.000	0.000	0.767	0.000
** INATS - INATS - Pilot Lot & Small Scale Manufacturing	C/CPFF	Southwest Research Institute San Antonio, TX	6.653	1.790	Jan 2010	0.000		0.000		0.000	0.000	8.443	0.000
INATS - Scale-Up Manufacturing of Candidate Oximes	C/CPIF	TBD	0.000	2.375	Apr 2010	0.000		0.000		0.000	0.000	2.375	0.000
Subtotal			6.653	4.932		0.000		0.000		0.000	0.000	11.585	0.000

Remarks

Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** INATS - INATS - Regulatory Integration, IND, and NDA Support Efforts	MIPR	Defense Technical Information Center Edgewood, MD	1.890	0.372	Jan 2010	0.000		0.000		0.000	0.000	2.262	0.000
Subtotal			1.890	0.372		0.000		0.000		0.000	0.000	2.262	0.000

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT MC4: <i>MEDICAL CHEMICAL DEFENSE (ACD&P)</i>
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Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			

Remarks

Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** IA - Inhalational Atropine - Formulation, Analytical Methods & Device Optimization	C/CPIF	TBD	0.000	1.200	Jan 2010	0.000		0.000		0.000	0.000	1.200	0.000
** INATS - INATS - Conduct Studies of Candidate Oximes	MIPR	Defense Technical Information Center Edgewood, MD	0.000	2.500	Apr 2010	0.000		0.000		0.000	0.000	2.500	0.000
Subtotal			0.000	3.700		0.000		0.000		0.000	0.000	3.700	0.000

Remarks

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT MC4: <i>MEDICAL CHEMICAL DEFENSE (ACD&P)</i>
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Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** INATS - INATS - Product Management Support	MIPR	USAMMDA Fort Detrick, MD	0.380	0.148	Jan 2010	0.000		0.000		0.000	0.000	0.528	0.000
INATS - Chem Bio Medical Systems	Allot	CBMS Frederick, MD	1.242	0.168	Apr 2010	0.000		0.000		0.000	0.000	1.410	0.000
** ZSBIR - SBIR/STTR - Aggregated from ZSBIR-SBIR/STTR	MIPR	HQ AMC, Alexandria	0.000	0.118		0.000		0.000		0.000	0.000	0.118	0.000
Subtotal			1.622	0.434		0.000		0.000		0.000	0.000	2.056	0.000

Remarks

	Total Prior Years Cost	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
Project Cost Totals	10.165	9.438	0.000	0.000	0.000	0.000	19.603	0.000

Remarks

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT MC4: <i>MEDICAL CHEMICAL DEFENSE (ACD&P)</i>
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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
** BSCAV - BSCAV Inc. 2 - Phase 1 Clinical Safety Studies	■	■	■	■																								
BSCAV Inc. 2 - Large Scale Manufacturing, Process Development & Assay Validation	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■				
BSCAV Inc. 2 - Milestone B					■																							
BSCAV Inc. 2 - Conduct GLP Animal Efficacy Studies							■	■	■	■	■	■	■	■	■	■	■	■	■	■								
BSCAV - NTA Studies	■	■	■	■																								
** IA - IA - Process Development and current Good Manufacturing Practices (cGMP) requirements							■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
IA - Formulation, analytical assay, and device development							■	■	■	■	■	■	■	■	■	■												
IA - Milestone B								■																				
** INATS - INATS - Process development & small scale cGMP	■	■	■	■	■	■	■	■																				
INATS - IND Application/Amendment	■	■	■	■	■	■	■	■																				
INATS - Efficacy, Safety & Toxicology Studies of Candidate Oximes		■	■	■	■	■	■	■																				
INATS - Process Development and small scale cGMP Manufacturing	■	■	■	■	■	■	■	■																				

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT MC4: <i>MEDICAL CHEMICAL DEFENSE (ACD&P)</i>
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Schedule Details

Event	Start		End	
	Quarter	Year	Quarter	Year
** BSCAV - BSCAV Inc. 2 - Phase 1 Clinical Safety Studies	4	2007	4	2009
BSCAV Inc. 2 - Large Scale Manufacturing, Process Development & Assay Validation	1	2008	4	2014
BSCAV Inc. 2 - Milestone B	2	2010	2	2010
BSCAV Inc. 2 - Conduct GLP Animal Efficacy Studies	3	2010	4	2013
BSCAV - NTA Studies	4	2008	4	2009
** IA - IA - Process Development and current Good Manufacturing Practices (cGMP) requirements	3	2010	3	2015
IA - Formulation, analytical assay, and device development	3	2010	4	2012
IA - Milestone B	4	2010	4	2010
** INATS - INATS - Process development & small scale cGMP	2	2008	3	2010
INATS - IND Application/Amendment	2	2008	4	2010
INATS - Efficacy, Safety & Toxicology Studies of Candidate Oximes	2	2009	4	2010
INATS - Process Development and small scale cGMP Manufacturing	1	2008	4	2010

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program								DATE: February 2010			
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>				R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>				PROJECT MR4: <i>MEDICAL RADIOLOGICAL DEFENSE (ACD&P)</i>			
COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
MR4: <i>MEDICAL RADIOLOGICAL DEFENSE (ACD&P)</i>	4.294	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	Continuing	Continuing
Quantity of RDT&E Articles											

A. Mission Description and Budget Item Justification

This project funds the advanced development of candidate therapeutic medical countermeasures to mitigate the consequences of exposure to ionizing radiation due to nuclear or radiological attacks. Exposure to ionizing radiation causes damage to blood-forming cells (hematopoietic system) and gastrointestinal system, leading to Acute Radiation Syndrome (ARS). Medical countermeasures must be approved by the Food and Drug Administration (FDA) for human use prior to fielding. Testing the efficacy of candidate drugs against normally lethal radiation exposure cannot be conducted in humans; therefore, surrogate animal models must be used to obtain FDA approval.

Medical Radiological Countermeasures (MRADC) efforts include multiple countermeasures required to restore casualties to pre-exposure health and to protect U.S. Forces against injury caused by exposure to radiation. MRADC shall reverse or limit radiation injury resulting in increased survival, decreased incapacity, and sustained operational effectiveness. In addition, MRADC shall be effective against a broad range of radiation sources and types and shall be useable in the battle space, including evacuation.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) MRADC <i>FY 2009 Accomplishments:</i> Completed initial non-clinical efficacy studies.	2.557	0.000	0.000	0.000	0.000
2) MRADC <i>FY 2009 Accomplishments:</i> Completed product formulation studies.	1.737	0.000	0.000	0.000	0.000

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT MR4: <i>MEDICAL RADIOLOGICAL DEFENSE (ACD&P)</i>
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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
Accomplishments/Planned Programs Subtotals	4.294	0.000	0.000	0.000	0.000

C. Other Program Funding Summary (\$ in Millions)

Line Item	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	FY 2012	FY 2013	FY 2014	FY 2015	Cost To Complete	Total Cost
• MR5: <i>MEDICAL RADIOLOGICAL DEFENSE (SDD)</i>	3.002	8.276	1.143		1.143	4.817	2.265	0.000	0.000	Continuing	Continuing

D. Acquisition Strategy

MRADC

Medical Identification and Treatment Systems (MITS) Joint Product Management Office will manage the development of Medical Radiation Countermeasures (MRADC) for the Department of Defense (DoD). A contractor will serve as the product integrator throughout development and shall be responsible for conducting activities associated with drug development in a manner consistent with eventual approval by the Food and Drug Administration (FDA). The contractor shall sponsor the drug to the FDA and hold all approvals and/or licenses. The Technology Development phase includes pre-clinical studies and Phase 1 human clinical safety studies. During the Engineering and Manufacturing Development (EMD) phase, large scale manufacturing, Phase 2 human clinical safety studies and definitive animal efficacy studies will be conducted. FDA approval of the countermeasure is an exit criterion for the EMD phase. During the Production and Deployment Phase, sufficient quantities of product to meet Initial Operational Capability and Full Operational Capability will be purchased. Subsequent purchases will be made by the Defense Logistics Agency. Any post-marketing surveillance requested by the FDA will be conducted.

MRADC will be developed using a system-of-systems approach to provide a full spectrum capability to protect against the radiation threat. Individual countermeasure solutions will be developed using a single step to a full capability (FDA approval) strategy. The DoD is working very closely with the Department of Health and Human Services (HHS), which also has a radiation countermeasure program. The establishment of an interagency working group provides oversight and guidance to both agency programs to ensure that their efforts are non-duplicative and are directed to meeting the requirement of both agencies.

E. Performance Metrics

N/A

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT MR4: <i>MEDICAL RADIOLOGICAL DEFENSE (ACD&P)</i>
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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
** MRADC - MRADC - Initial Non-Clinical Animal Efficacy Studies for 2 Candidates	■	■	■	■																								
MRADC - Product Formulation on Candidate 1	■	■	■	■																								
MRADC - Milestone B							■																					

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT MR4: <i>MEDICAL RADIOLOGICAL DEFENSE (ACD&P)</i>
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Schedule Details

Event	Start		End	
	Quarter	Year	Quarter	Year
** MRADC - MRADC - Initial Non-Clinical Animal Efficacy Studies for 2 Candidates	2	2008	4	2009
MRADC - Product Formulation on Candidate 1	2	2008	4	2009
MRADC - Milestone B	3	2010	3	2010

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program								DATE: February 2010			
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>				R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>				PROJECT TE4: <i>TEST & EVALUATION (ACD&P)</i>			
COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
TE4: <i>TEST & EVALUATION (ACD&P)</i>	6.261	28.773	19.304	0.000	19.304	11.851	28.035	20.266	21.139	Continuing	Continuing
Quantity of RDT&E Articles											

A. Mission Description and Budget Item Justification

This funding supports the Joint Project Manager Nuclear, Biological, Chemical Contamination Avoidance Product Director, Test Equipment, Strategy, and Support (PD TESS) efforts. PD TESS provides test infrastructure products for testing and evaluating chemical and biological defense systems throughout the life cycle acquisition process in support of the Milestone Decision Authority, Joint Project Managers, and the Test and Evaluation (T&E) community. PD TESS test infrastructure products are aligned in four groups to include: (1) Chemical Laboratory (Sense); (2) Biological Laboratory (Sense); (3) Individual Protection, Collective Protection and Decontamination (Shield and Sustain); and (4) Field Simulant (Sense).

(1) Chemical Laboratory (Sense): Products for this area include a Non-Traditional Agent (NTA) Test Facility. The NTA Facility provides a new capability at the Edgewood Chemical Biological Center (ECBC) to conduct highly toxic materials testing using new, emerging threat agents. The NTA facility supports testing of decontamination, collective protection, individual protection, and contamination avoidance products. Major CBDP acquisition programs supported are: the Joint Chemical Agent Detector (JCAD); the Joint Service General Purpose Mask (JSGPM); the Lightweight CB Ensemble (LCBE); Joint Expeditionary Collective Protection (JECBP); Joint Collective Protection Equipment (JCPE); Joint Service Transportable Decontamination System (JSTDS); Joint Warning and Reporting Network (JWARN) hardware components; the Joint Protective Aircrew Ensemble (JPACE); the Joint Service Aircrew Mask (JSAM); and the Joint Chemical Ensemble (JCE).

(2) Sense Laboratory (Biological): Products for this area include a biological live agent standoff chamber. The Chamber supports Joint Biological standoff detection testing in biological live agent environments. Major CBDP acquisition program support is the Joint Biological Standoff Detection System (JBSDS) Block II.

(3) Individual Protection, Collective Protection and Decontamination (Shield and Sustain): Products for this area include: Chemical, Biological Agent Resistance Test (CBART) fixture and Decontamination Coupon Validation Effort. CBART fixture provides a near real time testing capability under a range of environmental conditions for individual and collective protection materials. The Decontamination Coupon Validation Effort develops laboratory procedures to reduce variability in decontamination coupon testing. Acquisition Programs supported are: Joint Platform Interior Decontamination/Joint Material Decontamination System (JPID/JMDS); Joint Service Transportable Decontamination System (JSTDS); Joint Expeditionary Collective Protection (JECBP); Joint Collective Protection Equipment (JCPE); Protective Clothing; Joint Protective Aircrew Ensemble (JPACE); Joint Service General Purpose Mask (JSGPM); Joint Service Aircrew Mask (JSAM); and the Joint Chemical Ensemble (JCE).

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT TE4: <i>TEST & EVALUATION (ACD&P)</i>				
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) PD TESS - Non-Traditional Agent (NTA) Test System <i>FY 2009 Accomplishments:</i> Evaluated/optimized mock-up performance. Initiated test fixtures design and procedures development. <i>FY 2010 Plans:</i> Complete NTA Test Facility final design and initiate fabrication. Continue procedure and test fixtures development. <i>FY 2011 Base Plans:</i> Continue facility fabrication.		4.915	22.908	16.000	0.000	16.000
2) PD TESS - Bio Standoff Facility <i>FY 2010 Plans:</i> Initiate and complete Bio Standoff Facility feasibility effort. <i>FY 2011 Base Plans:</i> Initiate Bio Standoff Facility design.		0.000	0.600	0.900	0.000	0.900
3) PD TESS - Chemical Biological Agent Resistance Test (CBART) <i>FY 2011 Base Plans:</i> Initiate design.		0.000	0.000	0.900	0.000	0.900
4) PD TESS		1.346	3.904	1.504	0.000	1.504

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT TE4: <i>TEST & EVALUATION (ACD&P)</i>
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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2009 Accomplishments:</i> Continued systems engineering support for integration and execution of Advanced Component Development & Prototype development. <i>FY 2010 Plans:</i> Continue systems engineering support. <i>FY 2011 Base Plans:</i> Continue systems engineering support.					
5) PD TESS - Decon Coupon Validation <i>FY 2010 Plans:</i> Complete lab procedures for reduced variability.	0.000	1.000	0.000	0.000	0.000
6) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.	0.000	0.361	0.000	0.000	0.000
Accomplishments/Planned Programs Subtotals	6.261	28.773	19.304	0.000	19.304

C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011 Base</u>	<u>FY 2011 OCO</u>	<u>FY 2011 Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To Complete</u>	<u>Total Cost</u>
• TE5: <i>TEST & EVALUATION (SDD)</i>	37.444	36.593	15.901		15.901	12.243	4.238	14.614	15.300	Continuing	Continuing
	7.037	4.870	4.813		4.813	4.779	4.750	5.660	5.615	Continuing	Continuing

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT TE4: <i>TEST & EVALUATION (ACD&P)</i>
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C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011</u> <u>Base</u>	<u>FY 2011</u> <u>OCO</u>	<u>FY 2011</u> <u>Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To</u> <u>Complete</u>	<u>Total Cost</u>
• TE7: <i>TEST & EVALUATION (OP SYS DEV)</i>											

D. Acquisition Strategy

PD TESS

The PD TESS program provides for the development and acquisition of new and enhanced test infrastructure to support the sense, shield, shape, and sustain mission areas for the Joint Service Chemical and Biological Defense Program (CBDP). The efforts are supported through competitive contract actions, academia, and other Government agencies. Infrastructure solutions will leverage commercially available systems to provide state-of-the-art capabilities that address current and future CBDP test and evaluation needs.

E. Performance Metrics

N/A

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT TE4: <i>TEST & EVALUATION (ACD&P)</i>
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Product Development (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total		Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost	Award Date			
** PD TESS - HW S - NTA Test Facility Procedures/Fixtures	MIPR	Various	8.505	5.190	Jan 2010	2.000	Jan 2011	0.000		2.000	0.000	15.695	0.000	
HW S - NTA Test System Design/ Fabrication/Installation	C/FFP	TBD	3.997	18.018	Jan 2010	14.900	Jan 2011	0.000		14.900	0.000	36.915	0.000	
HW S - Bio Standoff Facility Feasibility/ Design	Reqn	TBD	0.000	0.600	Jan 2010	0.900	Jan 2011	0.000		0.900	0.000	1.500	0.000	
HW S - CBART - Design	C/CPPF	TBD	0.000	0.700	Apr 2010	0.000		0.000		0.000	0.000	0.700	0.000	
Subtotal			12.502	24.508		17.800		0.000		17.800	0.000	54.810	0.000	

Remarks

Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total		Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost	Award Date			
** PD TESS - PM/MS S - Program Management/ Systems Engineering Support	MIPR	JPM NBC CA APG, MD	4.694	3.904	Oct 2009	1.504	Oct 2010	0.000		1.504	0.000	10.102	0.000	
	MIPR	HQ	0.000	0.361		0.000		0.000		0.000	0.000	0.361	0.000	

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT TE4: <i>TEST & EVALUATION (ACD&P)</i>
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Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** ZSBIR - SBIR/STTR - Aggregated from ZSBIR-SBIR/STTR		AMC, Alexandria											
Subtotal			4.694	4.265		1.504		0.000		1.504	0.000	10.463	0.000

Remarks

	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract		
		Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost					
Project Cost Totals		17.196	28.773			19.304		0.000		19.304	0.000	65.273	0.000

Remarks

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT TE4: <i>TEST & EVALUATION (ACD&P)</i>
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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
** PD TESS - Bio Standoff Facility Design/ Fabrication/Installation						■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
DECON Coupon Validation					■	■	■	■																				

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT TE4: <i>TEST & EVALUATION (ACD&P)</i>
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Schedule Details

Event	Start		End	
	Quarter	Year	Quarter	Year
** PD TESS - Bio Standoff Facility Design/Fabrication/Installation	2	2010	4	2015
DECON Coupon Validation	1	2010	4	2010

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program									DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>				R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>				PROJECT TT4: <i>TECHBASE TECHNOLOGY TRANSITION (ACD&P)</i>			
COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
TT4: <i>TECHBASE TECHNOLOGY TRANSITION (ACD&P)</i>	17.065	26.649	26.466	0.000	26.466	18.564	18.838	19.294	19.563	Continuing	Continuing
Quantity of RDT&E Articles											

A. Mission Description and Budget Item Justification

This project (TT4) validates high-risk/high-payoff technologies, concepts-of-operations, and reconnaissance and surveillance platforms that could significantly improve Warfighter capabilities in preparation for transition of mature technologies to advanced development programs requiring chemical and biological (CB) defense technologies. These programs offer an opportunity to identify and efficiently mature emerging technologies from laboratory experiments to acquisition programs through risk reduction, engineering and integration. These Advanced Technology Demonstrations (ATDs) and Advanced Concept Technology Demonstrations (ACTDs) seek to demonstrate the potential for enhanced military operational capability and/or cost effectiveness. Upon conclusion of the technical and operational demonstrations, the user or sponsor provides a determination of the military utility and operational impact of the technology and capability demonstrated. Successfully demonstrated technologies with proven military utility can either be left in place for extended user evaluations, accepted into advanced stages of the formal acquisition process, proceed directly into limited or full-scale production or be returned to the technical base for further development. This project funds three major thrust areas (two of which are new thrust areas that consolidate legacy systems and are annotated as such below): Advanced Remediation Technologies (ART), Early Warning Military Application in Reconnaissance Systems (EW-MARS), and Comprehensive Innovative Protection (CIP). The ART addresses Chemical, Biological, and Radiological (CBR) remediation and decontamination processes and demonstrates technologies and methods to restore assets such as mobile equipment, fixed sites, critical infrastructures, personal, and equipment to operational status as a result of having reduced or eliminated CBR contamination. The EW-MARS (new thrust area) achieves enhanced command and control decision making capabilities as a result of a combined and orchestrated family of chemical and biological defense systems deployed on various platforms in remote locations. The CIP (new thrust area) transitions mature technologies to improve individual and collective protection capabilities for U.S. and coalition Warfighters. The following is a description of specific efforts funded under each thrust area:

ART:

Interagency Biological Restoration Demonstration (IBRD) - A Department of Defense (DoD)/Department of Homeland Security (DHS) collaborative effort that will provide a coordinated, systems approach to the recovery and restoration of wide urban areas. This will include Department of Defense (DoD) infrastructures and high traffic areas (transit/transportation facilities) following the aerosol release of a biological agent.

Special Platform Interior Decontamination and Equipment Restoration (SPIDER) - A concept exploration effort that focused on decontaminating the interior of an aircraft following the release of a chemical agent. This effort merged into the Decontamination Family of Systems, also known as HaMMER (see below for description).

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program	DATE: February 2010
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APPROPRIATION/BUDGET ACTIVITY	R-1 ITEM NOMENCLATURE	PROJECT
0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	TT4: <i>TECHBASE TECHNOLOGY TRANSITION (ACD&P)</i>

Automated Detailed Equipment Decontamination for Land Vehicles (Auto Decon) - A chemical and biological decontamination process for land vehicles, which will prototype an improved decontamination process and will evaluate the current Detailed Equipment Decontamination (DED), which is the most thorough of Joint Service decontamination procedures. This effort will merge into the Decontamination Family of Systems, also known as HaMMER (see below for description).

Hazard Mitigation Material and Equipment Restoration (HaMMER) - A layered strategy to identify individual technologies that may be collectively applied to reduce or eliminate chemical and biological hazards. It includes a Decontamination Family of Systems that gives the Warfighter multiple capabilities to reduce or eliminate chemical hazards. This effort leverages upon and consolidates Auto Decon and SPIDER efforts described above.

EW-MARS:

Military Applications in Reconnaissance Systems for Joint Force Protection (MARS-JFP) - A data fusion ATD that leverages early warning technologies developed in Budget Activity 3 (Project TT3) to improve the capability to detect and react to an initial chemical and biological attack, as well as prevent a second attack. Specifically, this effort focuses on force protection decision making for external, cross domain sensors for cueing/tipping, and managing resources of dynamically deployable high quality chemical and biological sensors.

Rapid Area Surveillance Reconnaissance (RASR) - A sensitive-site exploration, standoff reconnaissance, ATD that leverages early warning technologies developed in Budget Activity 3 (Project TT3) to survey large areas (whole rooms, courtyards, fields) and assess and identify contamination with Chemical Warfare Agents (CWAs), Toxic Industrial Chemicals (TICs) and Non-Traditional Agents (NTAs).

Post Intercept Weapons of Mass Destruction Identification (PIWID) - An ATD that leverages early warning technologies developed in Budget Activity 3 (Project TT3), which addresses both operational and technical issues associated with the capability to determine the presence of Weapons of Mass Destruction (WMD) in the threat payload of ballistic or cruise missile delivery systems after a successful active defense intercept.

CIP:

Demo-Low Burden Individual Protection Demonstration (IP Demo) - An ATD that leverages lightweight chemical and biological protective textiles developed in Budget Activity 3 (Project CB3, Protection and Hazard Mitigation), and will support the next generation Joint Chemical Ensemble. This effort will provide significantly decreased thermal burden correlated with acceptable levels of chemical and biological protection, as well as significantly increase the ability of the Warfighter to accomplish a mission in a contaminated environment.

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT TT4: <i>TECHBASE TECHNOLOGY TRANSITION (ACD&P)</i>				
Joint Medical Distance Support and Evaluation (JMDSE) - A Joint Concept Technology Demonstration (JCTD) that leverages the results of the EBD (see above for description) and seeks new detect-to-treat concept of operations (CONOPS) enabled by the deployment of new chemical and biological detection and identification capabilities to front line forces.						
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) TT DEMO ART (Interagency Biological Restoration Demonstration (IBRD)): <i>FY 2009 Accomplishments:</i> Continued development of restoration plans. Continued risk assessment. Conducted decontamination technologies efficacy testing relevant to an outdoor urban environment. Conducted agent fate and transport studies and demonstrations. Continued development and demonstration of system tools. Conducted table top exercises, field exercises, and workshops. <i>FY 2010 Plans:</i> Complete IBRD development of restoration plans. Complete established risk assessment and clearance goals. Develop sampling, characterization, and long term monitoring plans. Develop and exercise wide-area decontamination methods. Develop and demonstrate restoration system tools and conduct table top exercises, field exercises, and workshops. Plan, coordinate, and execute the IBRD Final Demo/Table Top Exercise (TTX) in the Seattle urban area. Transition decontamination methods, restoration tools, agent fate and transport data to the advanced developer (Joint Program Manager for Guardian and Decontamination - see Budget Activities 4 and 5).		5.827	2.716	0.000	0.000	0.000
2) TT DEMO ART (Automated Detailed Equipment Decontamination for Land Vehicles (Auto Decon)): <i>FY 2009 Accomplishments:</i> Initiated and conducted Test and Evaluation (T&E) of current detailed equipment decontamination processes and prototype automated decontamination solutions.		3.000	2.951	0.000	0.000	0.000

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT TT4: <i>TECHBASE TECHNOLOGY TRANSITION (ACD&P)</i>				
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2011 Base Plans:</i> Continue operational concept planning and exercise planning; technology readiness assessments; initiate operational mockup, lesson plans and final development planning; downselect prototype industry awards; conduct and finalize surety testing; conduct several technical and operational demonstrations; conduct Military Utility Assessment (MUA) to assess value to Warfighter; recondition complete systems in preparation for transition to operational managers and combat developers.						
6) TT DEMO EW-MARS Thrust Area (Post Intercept Weapons of Mass Destruction Identification (PIWID)): <i>FY 2009 Accomplishments:</i> Initiated Joint Land Attack Cruise Missile Elevated Netted Sensor (JLENS) study. Leveraged a missile intercept event for information gathering and baseline study. Conducted table top exercise to evaluate current Techniques, Tactics, and Procedures (TTPs). <i>FY 2010 Plans:</i> Conduct post-intercept WMD simulant payload data collection while leveraging missile intercept event. Demonstrate sidecar re-processing of non-chemical and biological sensors to extract useful cue/tipping information. <i>FY 2011 Base Plans:</i> Assess standoff data, chem/bio data, and current plan for Unmanned Aerial Vehicle (UAV) point-based, sensor approaches. Conduct standoff sensor and UAV CONOPS. Laboratory demonstration within cross domain environment. Transition data to JPM-NBC CA and JPM-BD.		2.000	1.968	1.796	0.000	1.796
7) TT DEMO CIP (Low Burden Individual Protection Demonstration (IP Demo)):		0.000	3.050	0.000	0.000	0.000

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT TT4: <i>TECHBASE TECHNOLOGY TRANSITION (ACD&P)</i>				
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2010 Plans:</i> Perform and complete system level technical performance measure evaluations. Initiate and complete a system level user demonstration. Conduct and complete component level testing. Transition low burden individual protection overgarment to the advanced developer (Joint Program Manager for Individual Protection and the Program Manager for Soldier Equipment). Initiate risk reduction activities to demonstrate Catalytic Oxidation air purification prototypes, to include initial system and tank test bed designs.						
8) TT DEMO CIP (Joint Medical Distance Support and Evaluation (JMSDE)): <i>FY 2009 Accomplishments:</i> Initiated internal planning, program management, and documentation. Conducted overlay scenarios and initiated JMDSE to Joint Biological Tactical Detection System (JBTDs) interface evaluation. <i>FY 2010 Plans:</i> Complete JMDSE to JBTDs interface evaluation. Conduct field demonstrations and military utility assessments. Develop CONOPS, training, test and security plans. Initiate software development. <i>FY 2011 Base Plans:</i> Complete field demonstrations and military utility assessments; complete CONOPS and training, test, and security plans. Complete software development and integration. Transition to JPM-Bio Detection.		0.738	0.984	1.505	0.000	1.505
9) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.		0.000	0.322	0.000	0.000	0.000
Accomplishments/Planned Programs Subtotals		17.065	26.649	26.466	0.000	26.466

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT TT4: <i>TECHBASE TECHNOLOGY TRANSITION (ACD&P)</i>
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C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011</u> <u>Base</u>	<u>FY 2011</u> <u>OCO</u>	<u>FY 2011</u> <u>Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To</u> <u>Complete</u>	<u>Total Cost</u>
• TE3: <i>TEST & EVALUATION (ATD)</i>	25.761	13.307	11.875		11.875	11.267	11.160	0.000	0.000	Continuing	Continuing
• TT3: <i>TECHBASE TECHNOLOGY TRANSITION</i>	8.127	7.357	4.504		4.504	8.117	8.169	8.390	8.528	Continuing	Continuing

D. Acquisition Strategy

TT DEMO

The Advanced Technology Demonstrations (ATDs) and Advanced Concept Technology Demonstrations (ACTDs) exploit mature and maturing technologies to solve important military problems. ATDs and ACTDs emphasize technology assessment and integration rather than technology development. The goal is to provide a prototype capability to the Warfighter and to support in the evaluation of that capability. The Warfighters evaluate the capabilities in real military exercises and at a scale sufficient to fully assess military utility. When possible, the ATDs will leverage results from existing chemical and biological science and technology (S&T) efforts and prior ATDs. Market research/baselining is performed prior to ATD initiation to determine if a suitable solution exists or whether a solicitation/sole source is required to develop a solution. The ATDs are typically managed by DoD, Federally Funded Research Development Centers (FFRDCs) or University Affiliated Research Centers (UARCs). This is done through the Military Interdepartmental Purchase Request (MIPR) or the Interagency Cost Reimbursable Order (IACRO) in accordance with the Economy Act. The ATD manager, who is typically responsible for total system development, can subcontract industry, academia, or other government agencies to perform individual component development.

E. Performance Metrics

N/A

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT TT4: <i>TECHBASE TECHNOLOGY TRANSITION (ACD&P)</i>
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Product Development (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** TT DEMO - HW S - (ART) IBRD System Design and Integration	PO	Pacific Northwest National Laboratory Seattle, WA	0.375	0.791	Jan 2010	0.000		0.000		0.000	0.187	1.353	0.000
HW C - (ART) Auto Decon	MIPR	Army- ECBC Edgewood, MD	0.000	0.300	Oct 2009	0.000		0.000		0.000	0.000	0.300	0.000
HW C - (ART) HaMMER Product Development	MIPR	Army- ECBC Edgewood, MD	0.000	2.950	Oct 2009	2.850	Oct 2010	0.000		2.850	0.000	5.800	0.000
HW S - (ART) Hammer Product Development-SME	MIPR	Army- ECBC Edgewood, MD	0.000	0.200	Oct 2009	0.200	Oct 2010	0.000		0.200	0.000	0.400	0.000
HW C - (EW) MARS JFP Product Development	PO	MITRE Bedford, MA	0.000	0.200	Jan 2010	0.200	Jan 2011	0.000		0.200	0.000	0.400	0.000
HW C - (EW) MARS JFP Product Development #2	PO	Johns Hopkins Univ/Applied Physics Lab (JHU-APL) Laurel, MD	0.000	0.200	Jan 2010	0.200	Jan 2011	0.000		0.200	0.000	0.400	0.000
HW C - (EW) MARS JFP Product Development #3	PO	MIT/Lincoln Labs Lexington, MA	0.000	0.200	Jan 2010	0.200	Jan 2011	0.000		0.200	0.000	0.400	0.000
HW C - (EW) RASR Product Development	PO	MIT/Lincoln Labs Lexington, MA	0.000	1.700	Jan 2010	1.650	Jan 2011	0.000		1.650	0.000	3.350	0.000
HW C - (EW) RASR Product Development #2	PO	Georgia Tech Institute of Technology Atlanta, GA	0.000	0.500	Jan 2010	0.500	Jan 2011	0.000		0.500	0.000	1.000	0.000

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT TT4: <i>TECHBASE TECHNOLOGY TRANSITION (ACD&P)</i>
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Product Development (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
HW C - (EW) RASR Product Development #3	PO	MITRE Bedford, MA	0.000	0.000		1.150	Jan 2011	0.000		1.150	0.000	1.150	0.000
HW C - (EW) RASR Product Development #4	PO	John Hopkins University/ Applied Physics Laboratory Laurel, MD	0.000	0.000		1.150	Jan 2011	0.000		1.150	0.000	1.150	0.000
HW C - (EW) RASR Product Development #5	PO	Kansas City Plant (DOE) Kansas City, MO	0.000	0.000		1.150	Jan 2011	0.000		1.150	0.000	1.150	0.000
HW C - (EW) RASR Product Development #6	PO	Naval Postgraduate School Monterey, CA	0.000	0.000		1.150	Jan 2011	0.000		1.150	0.000	1.150	0.000
HW C - (EW) PIWID Product Development	MIPR	JLENS Huntsville, AL	0.000	0.500	Oct 2009	0.500	Oct 2010	0.000		0.500	0.000	1.000	0.000
HW C - (CIP) IP Demo Product Development	MIPR	US Army Natick Soldier RD&E Center Natick, MA	0.000	0.300	Oct 2009	0.200	Oct 2010	0.000		0.200	0.000	0.500	0.000
HW C - (CIP) JMDSE Product Development	MIPR	US Army Natick Soldier RD&E Center Natick, MA	0.000	0.150	Oct 2009	0.150	Oct 2010	0.000		0.150	0.000	0.300	0.000
Subtotal			0.375	7.991		11.250		0.000		11.250	0.187	19.803	0.000

Remarks

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT TT4: <i>TECHBASE TECHNOLOGY TRANSITION (ACD&P)</i>
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Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** TT DEMO - ILS C - (ART) IBRD TTP and CONOPS Development	MIPR	SPAWAR San Diego, CA	1.028	0.250	Oct 2009	0.000		0.000		0.000	0.371	1.649	0.000
ILS C - (ART) Auto Decon Support	MIPR	Edgewood Chemical and Biological Center (ECBC) Edgewood, MD	0.000	0.800	Oct 2009	0.000		0.000		0.000	0.000	0.800	0.000
ILS S - (ART) HaMMER System Support	MIPR	Research Development & Engineering Cmd (RDECOM), Edgewood	0.000	1.400	Oct 2009	1.414	Oct 2010	0.000		1.414	0.000	2.814	0.000
ILS S - (ART) Hammer OM Support	MIPR	US European Command (USEUCOM) Stuttgart, GE	0.000	0.150	Oct 2009	0.150	Oct 2010	0.000		0.150	0.000	0.300	0.000
ILS S - (ART) HaMMER Support	MIPR	Edgewood Chemical and Biological Center Edgewood, MD	0.000	0.500	Oct 2009	0.500	Oct 2010	0.000		0.500	0.000	1.000	0.000
ILS C - (EW) MARS JFP Support	MIPR	Edgewood Chemical and Biological Center Edgewood, MD	0.000	0.465	Oct 2009	0.965	Oct 2010	0.000		0.965	0.000	1.430	0.000
ILS C - (EW) RASR OM Support	MIPR	20th Support Command	0.000	0.215	Oct 2009	0.215	Oct 2010	0.000		0.215	0.000	0.430	0.000

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT TT4: <i>TECHBASE TECHNOLOGY TRANSITION (ACD&P)</i>
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Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
		Aberdeen Proving Ground, MD											
ILS C - (EW) RASR OM Support #2	MIPR	MARFORPAC (PACOM) Camp Smith, HI	0.000	0.220	Oct 2009	0.220	Oct 2010	0.000		0.220	0.000	0.440	0.000
ILS C - (EW) PIWID Support-Data Analysis	MIPR	Air Force Research Laboratory Wright Patterson AFB, OH	0.000	0.200	Oct 2009	0.200	Oct 2010	0.000		0.200	0.000	0.400	0.000
ILS C - (EW) PIWID Support-Data Analysis #2	MIPR	JLENS Huntsville, AL	0.000	0.200	Oct 2009	0.200	Oct 2010	0.000		0.200	0.000	0.400	0.000
ILS C - (CIP) IP Demo Component Support	MIPR	US Army Natick Soldier RD&E Center Natick, MA	0.000	0.275	Oct 2009	0.110	Oct 2010	0.000		0.110	0.000	0.385	0.000
ILS C - (CIP) JMDSE Support	MIPR	US Army Natick Soldier RD&E Center Natick, MA	0.000	0.200	Oct 2009	0.200	Oct 2010	0.000		0.200	0.000	0.400	0.000
Subtotal			1.028	4.875		4.174		0.000		4.174	0.371	10.448	0.000

Remarks

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT TT4: <i>TECHBASE TECHNOLOGY TRANSITION (ACD&P)</i>
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Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** TT DEMO - OTE C - (ART) IBRD Operational Test	Allot	DTRA Test and Evaluation (DTRA CXT) Albuquerque, NM	1.500	0.809	Oct 2009	0.000		0.000		0.000	0.590	2.899	0.000
OTE S - (ART) Auto Decon System Testing	MIPR	Army- ECBC Edgewood, MD	0.000	0.800	Jan 2010	0.000		0.000		0.000	0.000	0.800	0.000
OTE S - (ART) HaMMER System Testing	MIPR	Army- ECBC Edgewood, MD	0.000	0.750	Oct 2009	0.750	Oct 2010	0.000		0.750	0.000	1.500	0.000
OTE S - (ART) HaMMER T&E Oversight	MIPR	Army- ECBC Edgewood, MD	0.000	0.400	Oct 2009	0.400	Oct 2010	0.000		0.400	0.000	0.800	0.000
OTE C - (EW) MARS JFP Support	MIPR	US Army Environmental Command (AEC) Aberdeen, MD	0.000	0.400	Oct 2009	0.200	Oct 2010	0.000		0.200	0.000	0.600	0.000
OTE C - (EW) MARS JFP Support #2	MIPR	Dugway Proving Ground (DPG) DPG, UT	0.000	0.500	Oct 2009	0.300	Oct 2010	0.000		0.300	0.000	0.800	0.000
OTE C - (EW) RASR Component Testing	MIPR	US Army Environmental Command (AEC) Aberdeen, MD	0.000	0.225	Oct 2009	0.675	Oct 2010	0.000		0.675	0.000	0.900	0.000
OTE C - (EW) RASR Component Testing #2	MIPR	DPG DPG, UT	0.000	0.225	Oct 2009	0.675	Oct 2010	0.000		0.675	0.000	0.900	0.000
OTE C - (EW) RASR Component Testing #3	MIPR	US Army Developmental Test Command	0.000	0.226	Oct 2009	0.729	Oct 2010	0.000		0.729	0.000	0.955	0.000

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT TT4: <i>TECHBASE TECHNOLOGY TRANSITION (ACD&P)</i>
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Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
		Aberdeen, MD											
OTE C - (EW) PIWID Component Testing	MIPR	DPG DPG, UT	0.000	0.400	Oct 2009	0.400	Oct 2010	0.000		0.400	0.000	0.800	0.000
OTE C - (EW) PIWID Component Testing #2	MIPR	JLENS Huntsville, AL	0.000	0.400	Oct 2009	0.400	Oct 2010	0.000		0.400	0.000	0.800	0.000
OTE C - (CIP) IP Demo T&E	MIPR	US Army Natick Soldier RD&E Center Natick, MA	0.000	1.745	Oct 2009	1.200	Oct 2010	0.000		1.200	0.000	2.945	0.000
OTE C - (CIP) JMDSE Demo and Evaluation	MIPR	US Army Natick Soldier RD&E Center Natick, MA	0.000	0.400	Oct 2009	0.400	Oct 2010	0.000		0.400	0.000	0.800	0.000
Subtotal			1.500	7.280		6.129		0.000		6.129	0.590	15.499	0.000

Remarks

Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
	MIPR	Space and Naval Warfare Systems	1.326	0.950	Oct 2009	0.000		0.000		0.000	0.469	2.745	0.000

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT TT4: <i>TECHBASE TECHNOLOGY TRANSITION (ACD&P)</i>
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Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** TT DEMO - PM/MS S - IBRD Program Management		Command (SPAWAR) San Diego, CA											
PM/MS S - Auto Decon Management Support	MIPR	Army - ECBC Edgewood, MD	0.000	0.950	Oct 2009	0.000		0.000		0.000	0.000	0.950	0.000
PM/MS S - HaMMER System Management	MIPR	Army - ECBC Edgewood, MD	0.000	0.700	Oct 2009	0.740	Oct 2010	0.000		0.740	0.000	1.440	0.000
PM/MS S - HaMMER System Program Management	MIPR	Army - ECBC Edgewood, MD	0.000	0.875	Oct 2009	0.940	Oct 2010	0.000		0.940	0.000	1.815	0.000
PM/MS S - MARS JFP Program Management	MIPR	Army - ECBC Edgewood, MD	0.000	0.950	Oct 2009	0.985	Oct 2010	0.000		0.985	0.000	1.935	0.000
PM/MS S - RASR Program Management	MIPR	Army - ECBC Edgewood, MD	0.000	0.675	Oct 2009	1.500	Oct 2010	0.000		1.500	0.000	2.175	0.000
PM/MS S - PIWID System Program Management	MIPR	JLENS Huntsville, AL	0.000	0.270	Oct 2009	0.300	Oct 2010	0.000		0.300	0.000	0.570	0.000
PM/MS C - IP Demo Program Management	MIPR	US Army Natick Soldier RD&E Center Natick, MA	0.000	0.580	Oct 2009	0.200	Oct 2010	0.000		0.200	0.000	0.780	0.000
PM/MS C - JMDSE Program Management	MIPR	US Army Natick Soldier RD&E Center Natick, MA	0.000	0.231	Oct 2009	0.248	Oct 2010	0.000		0.248	0.000	0.479	0.000
	MIPR	HQ	0.000	0.322		0.000		0.000		0.000	0.000	0.322	0.000

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT TT4: <i>TECHBASE TECHNOLOGY TRANSITION (ACD&P)</i>
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Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** ZSBIR - SBIR/STTR - Aggregated from ZSBIR-SBIR/STTR		AMC, Alexandria											
Subtotal			1.326	6.503		4.913		0.000		4.913	0.469	13.211	0.000

Remarks
Management service costs cover all ten ATDs described in the R2a of this project (TT4).

	Total Prior Years Cost	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
Project Cost Totals	4.229	26.649	26.466	0.000	26.466	1.617	58.961	0.000

Remarks

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT TT4: <i>TECHBASE TECHNOLOGY TRANSITION (ACD&P)</i>
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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
** TT DEMO - (ART) Interagency Biological Restoration Demonstration (IBRD)	■	■	■	■	■	■	■	■																				
(ART) Automated Detailed Equipment Decontamination for Land Vehicles (Auto Decon)	■	■	■	■	■	■	■	■																				
(ART) Hazard Mitigation, Material and Equipment Restoration (HaMMER)	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■												
(EW) Military Applications in Reconnaissance/Support (MARS JFP)	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■								
(EW) Rapid Area-Scan Sensitive-site Reconnaissance (RASR)	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■												
(EW) Post Intercept WMD Identification (PIWID)	■	■	■	■	■	■	■	■	■	■	■	■																
(CIP) IP Demo	■	■	■	■	■	■	■	■	■	■	■	■																
(CIP) JMDSE	■	■	■	■	■	■	■	■	■	■	■	■																

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 4: <i>Advanced Component Development & Prototypes (ACD&P)</i>	R-1 ITEM NOMENCLATURE PE 0603884BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (ACD&P)</i>	PROJECT TT4: <i>TECHBASE TECHNOLOGY TRANSITION (ACD&P)</i>
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Schedule Details

Event	Start		End	
	Quarter	Year	Quarter	Year
** TT DEMO - (ART) Interagency Biological Restoration Demonstration (IBRD)	1	2007	4	2010
(ART) Automated Detailed Equipment Decontamination for Land Vehicles (Auto Decon)	1	2008	4	2010
(ART) Hazard Mitigation, Material and Equipment Restoration (HaMMER)	1	2009	4	2012
(EW) Military Applications in Reconnaissance/Support (MARS JFP)	1	2009	4	2013
(EW) Rapid Area-Scan Sensitive-site Reconnaissance (RASR)	1	2009	4	2012
(EW) Post Intercept WMD Identification (PIWID)	1	2009	4	2011
(CIP) IP Demo	1	2009	4	2011
(CIP) JMDSE	1	2009	4	2011

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Exhibit R-2, RDT&E Budget Item Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>
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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
Total Program Element	286.529	300.317	407.162	0.000	407.162	413.610	368.621	308.708	228.651	Continuing	Continuing
CA5: <i>CONTAMINATION AVOIDANCE (SDD)</i>	46.316	78.042	124.936	0.000	124.936	117.729	110.250	89.493	58.830	Continuing	Continuing
CM5: <i>HOMELAND DEFENSE (SDD)</i>	3.897	8.638	1.166	0.000	1.166	3.822	0.000	2.361	2.413	Continuing	Continuing
CO5: <i>COLLECTIVE PROTECTION (SDD)</i>	13.323	12.002	18.459	0.000	18.459	11.671	10.267	7.835	0.000	Continuing	Continuing
DE5: <i>DECONTAMINATION SYSTEMS (SDD)</i>	16.611	36.786	28.499	0.000	28.499	23.944	25.770	14.701	5.928	Continuing	Continuing
IP5: <i>INDIVIDUAL PROTECTION (SDD)</i>	18.363	21.094	9.678	0.000	9.678	4.833	3.044	0.756	0.563	Continuing	Continuing
IS5: <i>INFORMATION SYSTEMS (SDD)</i>	45.694	27.301	13.844	0.000	13.844	24.984	24.872	25.345	25.775	Continuing	Continuing
MB5: <i>MEDICAL BIOLOGICAL DEFENSE (SDD)</i>	87.676	57.558	141.680	0.000	141.680	161.732	159.144	141.481	111.671	Continuing	Continuing
MC5: <i>MEDICAL CHEMICAL DEFENSE (SDD)</i>	14.203	14.027	51.856	0.000	51.856	47.835	28.771	12.122	8.171	Continuing	Continuing
MR5: <i>MEDICAL RADIOLOGICAL DEFENSE (SDD)</i>	3.002	8.276	1.143	0.000	1.143	4.817	2.265	0.000	0.000	Continuing	Continuing
TE5: <i>TEST & EVALUATION (SDD)</i>	37.444	36.593	15.901	0.000	15.901	12.243	4.238	14.614	15.300	Continuing	Continuing

A. Mission Description and Budget Item Justification

Operational forces have an immediate need to survive, safely operate, and sustain operations in a chemical and biological agent threat environment across the continuum of global, contingency, special operations/low-intensity conflict, counter-narcotics, and other high risk missions. Operating forces have a critical need for defense against worldwide proliferation of Chemical and Biological (CB) warfare capabilities and for medical treatment of casualties in medical treatment facilities.

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Exhibit R-2, RDT&E Budget Item Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>
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Congress has directed centralized management of Department of Defense (DoD) CB Defense initiatives, both medical and non-medical. This program element supports the System Development and Demonstration (SDD) of CB defensive equipment, both medical and non-medical. These projects have been restructured to consolidate Joint and Service-unique tasks within four commodity areas: contamination avoidance, force protection (individual and collective), decontamination, and medical countermeasures. The consolidation will provide for development and operational testing of equipment for Joint Service as well as Service-unique requirements.

Contamination avoidance efforts under this system development program will provide U.S. forces with real-time hazard assessment capabilities. They include advanced multi-agent point and remote chemical detection systems for ground, aircraft, and shipboard applications; automated warning and reporting systems; integrated radiation detection and monitoring equipment; and enhanced battlefield reconnaissance capabilities. Force protection efforts will increase protection levels while decreasing physical and psychological burdens imposed by protective equipment. They include improved aircrew respiratory protection, lightweight integrated suit technology, and shipboard collective protection equipment.

Weapons of Mass Destruction Civil Support Team (WMD CST) efforts provide for testing and development of a Unified Command Suite (UCS) and an Analytical Laboratory Platform (ALS) for these teams.

The medical chemical defense system development program funds improved medical equipment and drugs essential to counteracting lethal and performance-degrading effects of chemical threats and medical equipment essential to meeting medical requirements on the integrated battlefield with emphasis on decreased size/weight and high mobility, yet supporting large numbers of combat casualties. Additionally, foreign medical materiel may be procured for exploitation of advanced technology and development to meet medical defense goals. This program element supports the development of prophylactic and therapeutic drugs and rapid identification and diagnostic systems. This program also funds development of a Transformational Rapid Drug Discovery and Development Capability (TRDDDC). Transformational Medical Technology Initiatives (TMTI) efforts in this area will include the continual build out of both a genomic sequencing and a bio-chemical informatics capability for the DoD.

DoD Biological Defense mission requires the detection of validated biological threat agents to provide early warning capabilities on mobile and fixed platforms. This program element will provide theater protection through the development of point and stand-off detection systems. The detection system concept will provide detection, identification, warning, and sample collection for verification that a biological agent attack has occurred. This program element also provides for the development of biological defense medical programs. DoD Biological Defense medical mission will address: (1) protective vaccines - vaccination capability against the most probable biological threat agents; (2) identification - clinical identification of biological threat agents through medical evaluation and laboratory analysis to augment early warning capabilities.

The projects in this program element support efforts in the system development phases of the acquisition strategy and are therefore correctly placed in Budget Activity 5.

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Exhibit R-2, RDT&E Budget Item Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY	R-1 ITEM NOMENCLATURE
0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i>	PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>
BA 5: <i>Development & Demonstration (SDD)</i>	

B. Program Change Summary (\$ in Millions)

	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011 Base</u>	<u>FY 2011 OCO</u>	<u>FY 2011 Total</u>
Previous President's Budget	300.149	332.895	0.000	0.000	0.000
Current President's Budget	286.529	300.317	407.162	0.000	407.162
Total Adjustments	-13.620	-32.578	407.162	0.000	407.162
• Congressional General Reductions		-48.658			
• Congressional Directed Reductions		0.000			
• Congressional Rescissions	0.000	0.000			
• Congressional Adds		16.080			
• Congressional Directed Transfers		0.000			
• Reprogrammings	-5.116	0.000			
• SBIR/STTR Transfer	-3.504	0.000			
• Other Adjustments	-5.000	0.000	407.162	0.000	407.162

Congressional Add Details (\$ in Millions, and Includes General Reductions)

Project: DE5: *DECONTAMINATION SYSTEMS (SDD)*

Congressional Add: 1) *Chemical and Biological Threat Reduction Coating*

Congressional Add: 2) *Self-Decontaminating Polymer System for Chemical and Biological Warfare Agents.*

Congressional Add: 3) *Self Contained Automated Vehicle Washing Systems with microwave decontamination*

Congressional Add: 4) *Protective Self-Decontaminating Surfaces*

Congressional Add Subtotals for Project: DE5

Project: IP5: *INDIVIDUAL PROTECTION (SDD)*

Congressional Add: 1) *JSAM*

Congressional Add Subtotals for Project: IP5

Project: TE5: *TEST & EVALUATION (SDD)*

Congressional Add: 1) *Real Time Monitoring of Chemical Agents*

	<u>FY 2009</u>	<u>FY 2010</u>
	0.000	2.390
	0.000	2.788
	0.000	1.593
	0.000	1.593
	0.000	8.364
	1.582	2.390
	1.582	2.390
	0.000	1.275

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Exhibit R-2, RDT&E Budget Item Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>
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Congressional Add Details (\$ in Millions, and Includes General Reductions)

	FY 2009	FY 2010
Congressional Add Subtotals for Project: TE5	0.000	1.275
Congressional Add Totals for all Projects	1.582	12.029

Change Summary Explanation

Funding: N/A - Adjustments less than 10% of total program.

Schedule: N/A

Technical: N/A

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program								DATE: February 2010			
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>				R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>				PROJECT CA5: <i>CONTAMINATION AVOIDANCE (SDD)</i>			
COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
CA5: <i>CONTAMINATION AVOIDANCE (SDD)</i>	46.316	78.042	124.936	0.000	124.936	117.729	110.250	89.493	58.830	Continuing	Continuing
Quantity of RDT&E Articles	89	0	0		0	0	0	0	0		

A. Mission Description and Budget Item Justification

This funding supports Engineering and Manufacturing Development and Low Rate Initial Production (EMD/LRIP) of an array of reconnaissance, detection and identification equipment, and warning systems.

Efforts funded in this project are: (1) Chemical, Biological, Radiological, and Nuclear Dismounted Reconnaissance Systems (CBRN DRS, formerly JNBCRS Increment 2); (2) Joint Biological Point Detection System (JBPDS); (3) Joint Biological Stand-off Detection System (JBSDS); (4) Joint Biological Tactical Detection System (JBTDS); (5) Joint Chemical Agent Detector (JCAD); (6) Joint Service Chemical Biological and Chemical Reconnaissance Systems Increments 2 and 3 (JNBCRS 2 and 3); (7) Major Defense Acquisition Program (MDAP) Support; (8) Next Generation Chemical Standoff Detection (NGCSD); (9) Non-Traditional Agent (NTA) Detection Support; and (10) Non-Traditional Agent Detection Sensor Suite Integration for NBC Reconnaissance Systems (SSI NBCRS) (formerly JNBCRS Increment 3).

The CBRN Dismounted Reconnaissance Systems (CBRN DRS) program fills a mission critical need to enhance CBRN reconnaissance platoon capabilities. The program consists of two phases. Phase I is the Dismounted Reconnaissance (DR) Sets, Kits and Outfits (SKO) configuration which fulfills an immediate critical need consisting of Commercial Off-The-Shelf (COTS) and Government Off-The-Shelf (GOTS) equipment integrated into a modular, transportable container for dismounted operations. It will form the basis for Phase II which is the Monitoring and Survey (MS) SKO. The MS SKO will feature technology insertion, the addition of net-centric capability and tailoring to focus on the Service-specific needs, to include NTA detection. The term "JNBCRS Increment 2" is replaced by the term "CBRN DRS" in FY10.

The JBPDS is a Joint Service biological detector system for the Services. The Army platforms include the JBPDS on the Biological Integrated Detection System (BIDS) and the Stryker Nuclear Biological Chemical Reconnaissance Vehicle (NBCRV). The Navy has identified select surface ships for installation of the JBPDS. The JBPDS is a fully automated system that increases the number of agents that can be identified by the current BIDS P3I and Interim Bio Agent Detector System (IBADS). Build 2, the JBPDS upgrade to Increment 1, will be developed. Build 2 will reduce lifecycle costs, improve reliability, and address system obsolescence concerns. The Build 2 program will incorporate one technology base transition of the Rapid Agent Aerosol Detector (RAAD) into a size, weight and power requirement to lower false alarms in the JBPDS which will help lower consumable use and reduce operations and support costs during its life cycle. Other JBPDS subsystem improvements are also focused on reductions to operational cost and obsolescence issues.

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
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<p>The JBSDS is employing an incremental acquisition strategy. JBSDS Increment 1 is the first standoff early warning biological detection (BD) system for the Joint Services. The system will be capable of providing near real time detection of biological attacks/incidents and standoff early detection/warning (Detect to Warn) of biological Warfare (BW) agents at fixed sites or in static mode on vehicles. It will be capable of providing standoff detection, ranging, tracking, discrimination (man-made vs. natural occurring aerosols) of BW aerosol clouds for advanced warning, reporting, and protection. The JBSDS will augment and integrate with existing BD systems to provide a BD network capable of near real time detection and warning theater-wide to limit the effects of biological agent hazards against U.S. forces at the tactical and operational levels of war. The JBSDS can be employed in support of various areas (e.g., fixed sites, Air Ports of Debarcation/Sea Ports of Debarcation (APODs/SPODs), amphibious landing sites, etc.), or on platforms (ships, aircraft or ground vehicles).</p> <p>The JBSDS Increment 2 builds on the capabilities demonstrated during the development of JBSDS Increment 1. The JBSDS Increment 2 system will focus on providing 24-hour operations (Increment 1 is night-time only), improving the false alarm rate and detection sensitivity, while decreasing size, weight and power. The JBSDS Increment 2 will also integrate with the global information network to provide near real time detection and warning theater-wide to limit the effect of biological agent hazards against U.S. forces at the tactical and operational levels of war. JBSDS Increment 3 will build on Increment 2 and focus on the development of a system that will operate on-the-move on mobile platforms as determined by the Warfighter.</p> <p>The Joint Biological Tactical Detection System (JBTDS) Increment 1 will develop, integrate, test and produce a lightweight, low cost biological surveillance system that will detect, collect and identify biological warfare agent aerosols. JBTDS will provide warning through the Joint Warning And Reporting Network (JWARN) and an archive sample for follow-on analyses. JBTDS will provide near-real-time local audio and visual alarm. JBTDS components will be man portable and battery operable. JBTDS will be used organically at Brigade and below and at Forward Operating Bases (FOB) to provide notification of a hazard and enhanced battle space awareness to protect and preserve the force. When networked, JBTDS will augment existing biological detection systems to provide a theater-wide seamless array capable of biological detection, identification, and warning.</p> <p>The JCAD program employs an incremental acquisition strategy to develop a miniaturized, rugged, and portable point chemical agent detector that automatically and simultaneously detects, identifies, quantifies, and alerts in the presence of nerve, blister, and blood chemical warfare agents. The M4 JCAD entered full rate production in September 2008 and will be produced through FY10. The attainable JCAD Increment 2 capabilities within the JCAD Increment 1 objectives were incorporated into a product improvement of the M4 JCAD (M4E1). Production of the M4E1 is scheduled to begin in FY11. JCAD will be used for wheeled vehicles, stand alone, and individual soldier applications. The M4 JCAD will replace the M8A1 and the M22 Automatic Chemical Agent Alarms (ACAA/ACADA). The M4E1 may additionally replace the Chemical Agent Monitor (CAM) and Improved Chemical Agent Monitor (ICAM) and other legacy systems currently used by the individual Services.</p> <p>The JNBCRS Increment 2 (which has been renamed to CBRN Dismounted Reconnaissance Systems for FY10) fills a mission critical need to enhance CBRN reconnaissance platoon capabilities. The program consists of two Phases. Phase I is the dismounted reconnaissance (DR) sets, kits and outfits (SKO) configuration which provides an immediate critical need consisting of COTS and GOTS integrated into a modular, transportable container for dismounted operations. It will form the</p>		

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<p>basis for Phase II which is the Monitoring and Survey (MS) SKO. JNBCRS Increment 2 initiated as a Joint Urgent Operational Needs Statement (JUONS) system to support the DR mission. The MS SKO will feature technology insertion, the addition of net-centric capability, and tailoring to focus on the service-specific needs, to include NTA detection.</p> <p>The JNBCRS Increment 3 will provide Chemical Biological Mass Spectrometer (CBMS) and Joint Contaminated Surface Detector (JCSD) capability to the Stryker Product Improvement Program and Future Mounted Armored Reconnaissance Platforms. The CBMS Bio effort will add the biological warfare agent detection and identification capability to the existing chemical liquid, and developmental Toxic Industrial Chemical (TIC) capabilities. The integration of liquid chemical and biological aerosol detection, within a single sensor; saves size, weight, and power on the platform. The JCSD will provide an improved mobile reconnaissance capability and on-the-move, non-contact, detection and identification of Chemical Warfare Agents (CWAs), TICs, and other Non-Traditional Agents (NTAs) using laser induced Raman Spectroscopy. Target surfaces are illuminated by laser light, and contaminants in the field of view are identified through analysis of their Raman backscatter signal against a wide library of Raman spectra. The JNBCRS Increment 3 supports transition to SSI NBCRS starting in FY10.</p> <p>The Major Defense Acquisition Program (MDAP) Support program will integrate System of Systems (SoS) solutions across the Armed Services for (MDAP) having Chemical and Biological Radiological and Nuclear (CBRN) survivability requirements. The program will demonstrate modular, net-centric, "plug and play" capabilities for mounted and dismounted CBRN reconnaissance that will establish a common CBRN reconnaissance architecture across the services.</p> <p>The Joint Science Lightweight Standoff Chemical Agent Detection (JSLSCAD) effort initiated the component improvements and the Technology Readiness Assessment (TRA) for the System of Systems (SoS) approach to address the CB early warning mission within the Next Generation Chemical Standoff Detection (NGCSD) program. The NGCSD SoS approach will increase the range of standoff detection and decrease detection time.</p> <p>The NGCSD effort will provide early warning for both traditional and non-traditional chemical agent attacks at fixed sites, forward operating bases and on Service designated vehicles and ships. This effort will develop and integrate new standoff sensor technologies for future standoff systems. The detection system will interoperate with the Services and Joint Command, Control, Communications, Computers, Intelligence, Surveillance and Reconnaissance (C4ISR) architectures.</p> <p>The Non-Traditional Agent (NTA) Detection program will develop and procure detection system(s) through incremental evolution that will afford Warfighter's the ability to attain situational awareness and respond to emerging hazards. The program will provide a near term capability to detect priority emerging threat materials with common core technologies for detection and identification. The common technologies can be further exploited in future increments to address lab deployable, fixed site and handheld applications.</p> <p>The SSI NBCRS will provide Chemical Biological Mass Spectrometer (CBMS) and Joint Contaminated Surface Detector (JCSD) capability to the Stryker Product Improvement Program and Future Mounted Armored Reconnaissance Platforms. The CBMS Bio effort will add the biological warfare agent detection and identification capability to the existing chemical liquid, and developmental Toxic Industrial Chemical (TIC) capabilities. The integration of liquid chemical and biological aerosol</p>		

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detection, within a single sensor; saves size, weight, and power on the platform. The JCSD will provide an improved mobile reconnaissance capability and on-the-move, non-contact, detection and identification of Chemical Warfare Agents (CWAs), TICs, and other Non-Traditional Agents (NTAs) using laser induced Raman Spectroscopy. Target surfaces are illuminated by laser light, and contaminants in the field of view are identified through analysis of their Raman backscatter signal against a wide library of Raman spectra. The SSI NBCRS was transitioned from JNBCRS Increment 3 in FY10.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) CBRN DRS <i>FY 2010 Plans:</i> Conduct engineering support (Govt). <i>FY 2011 Base Plans:</i> Continue engineering support (Govt).	0.000	0.718	1.407	0.000	1.407
2) CBRN DRS <i>FY 2010 Plans:</i> Initiate Developmental Testing to support Low Rate Initial Production (LRIP) decision. <i>FY 2011 Base Plans:</i> Complete Developmental Testing to support LRIP decision.	0.000	2.819	2.696	0.000	2.696
3) CBRN DRS <i>FY 2010 Plans:</i> Initiate Operational Assessment. <i>FY 2011 Base Plans:</i> Complete Operational Assessment.	0.000	2.300	2.800	0.000	2.800
4) CBRN DRS	0.000	2.400	6.600	0.000	6.600

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2009 Accomplishments:</i> Continued Modeling and Simulation.						
16) JBSDS Increment 2 <i>FY 2009 Accomplishments:</i> Conducted Agent Performance Assessment.		2.186	0.000	0.000	0.000	0.000
17) JBSDS Increment 1 <i>FY 2009 Accomplishments:</i> Provided Test Support.		0.493	0.000	0.000	0.000	0.000
18) JBSDS Increment 2 <i>FY 2009 Accomplishments:</i> Developed Test Equipment for Technology Demonstration V.		1.056	0.000	0.000	0.000	0.000
19) JBTDS <i>FY 2009 Accomplishments:</i> Initiated strategic/tactical planning, government systems engineering, program/financial management, costing, technology assessment, contracting, scheduling, acquisition oversight and technical support.		0.262	0.000	0.000	0.000	0.000
20) JCAD <i>FY 2009 Accomplishments:</i> Purchased M4E1 JCAD systems (Quantity of 89 at \$6,820 each for a total cost of \$707,000).		0.607	0.000	0.000	0.000	0.000
21) JCAD		5.631	3.360	0.000	0.000	0.000

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2009 Accomplishments:</i> Conducted M4E1 JCAD Developmental Tests to include surety chamber (\$3.2 million) and chamber upgrade testing (\$782 thousand).</p> <p><i>FY 2010 Plans:</i> Conduct M4E1 JCAD Developmental Tests to include surety chamber (\$2.175 million), MIL-STD-810/EMI (\$490 thousand), and false alarm and other testing (\$695 thousand).</p>						
<p>22) JCAD</p> <p><i>FY 2009 Accomplishments:</i> Provided Program Management Support.</p> <p><i>FY 2010 Plans:</i> Provide Systems Engineering (\$1.556 million), Program Management (\$950 thousand), and T&E IPT Support (\$700 thousand).</p> <p><i>FY 2011 Base Plans:</i> Provide Program Management and Systems Engineering Support.</p>		1.062	3.206	1.733	0.000	1.733
<p>23) JCAD</p> <p><i>FY 2010 Plans:</i> Conduct M4E1 JCAD Operational Test and Evaluation.</p>		0.000	1.510	0.000	0.000	0.000
<p>24) JCAD</p> <p><i>FY 2011 Base Plans:</i> Conduct M4E1 JCAD Multi-Service Operational Test and Evaluation (MOT&E).</p>		0.000	0.000	8.000	0.000	8.000
<p>25) JNBCRS INC 2</p>		9.794	0.000	0.000	0.000	0.000

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2009 Accomplishments:</i> Continued development of Joint Urgent Operational Needs (JUONS) and initiated development of program documentation, award contract for Integrated Logistics Support (ILS), design and test of DR-SKO.						
26) JNBCRS INC 2 <i>FY 2009 Accomplishments:</i> Continued DT/OT planning and other test agency support for JUONS.		2.500	0.000	0.000	0.000	0.000
27) JNBCRS INC 2 <i>FY 2009 Accomplishments:</i> Continued Systems Engineering Support (Govt) for JUONS and DR-SKO.		0.459	0.000	0.000	0.000	0.000
28) JNBCRS INC 3 <i>FY 2009 Accomplishments:</i> Completed design and development testing of Chemical Biological sensors.		1.578	0.000	0.000	0.000	0.000
29) JNBCRS INC 3 <i>FY 2009 Accomplishments:</i> Completed development of Common CBRN Sensor Interface (CCSI) prototype detectors.		0.436	0.000	0.000	0.000	0.000
30) JNBCRS INC 3 <i>FY 2009 Accomplishments:</i> Completed design of CCSI prototype detector housing (e.g. cradle).		0.052	0.000	0.000	0.000	0.000
31) JNBCRS 3		0.433	0.000	0.000	0.000	0.000

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2009 Accomplishments:</i> Completed engineering support.						
32) JSLSCAD <i>FY 2009 Accomplishments:</i> Provided Test Site Support for Next Generation Chemical Standoff Detection (NGCSD).		0.485	0.000	0.000	0.000	0.000
33) JSLSCAD <i>FY 2009 Accomplishments:</i> Conducted Sensor Hardware Development to Support the NGCSD Operational Demonstration.		0.245	0.000	0.000	0.000	0.000
34) JSLSCAD <i>FY 2009 Accomplishments:</i> Conducted Integrated Sensor Development and Testing from multiple vendors to support the NGCSD Operational Demonstration.		1.085	0.000	0.000	0.000	0.000
35) MDAP SPRT <i>FY 2009 Accomplishments:</i> Continued analysis and development of SoS architecture that supports MDAP operational CBRN requirements and provides Chemical Biological Radiological Nuclear (CBRN) defense capabilities.		2.125	0.000	0.000	0.000	0.000
36) MDAP SPRT <i>FY 2009 Accomplishments:</i> Continued Developmental Test (DT) to validate and verify SoS concept prior to MDAP integration.		3.410	0.000	0.000	0.000	0.000
37) MDAP SPRT		0.701	3.095	2.155	0.000	2.155

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
44) NGCSD <i>FY 2011 Base Plans:</i> Initiate engineering support for development of NGCSD.		0.000	0.000	1.500	0.000	1.500
45) NGCSD <i>FY 2011 Base Plans:</i> Initiate logistics planning efforts for manuals, maintenance, sparing, etc.		0.000	0.000	0.600	0.000	0.600
46) NGCSD <i>FY 2010 Plans:</i> Initiate JPEO-CBD Integrated Program Assistance Team (IPAT) support for Material Development Decision (MDD). <i>FY 2011 Base Plans:</i> Continue JPEO-CBD Integrated Program Assistance Team (IPAT) support for Material Development Decision (MDD).		0.000	1.500	0.750	0.000	0.750
47) NGCSD <i>FY 2010 Plans:</i> Initiate and complete test methodology development.		0.000	3.615	0.000	0.000	0.000
48) NTA DETECT <i>FY 2010 Plans:</i> Initiate Commercial Off-the-Shelf (COTS)/Government Off-the-Shelf (GOTS) evaluation for Installation Force Protection Mission Areas.		0.000	1.350	2.912	0.000	2.912

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2010 Plans:</i> (CBMS) Initiate engineering support, leveraging efforts initiated under Joint NBC Reconnaissance System (JNBCRS) INC 3. <i>FY 2011 Base Plans:</i> (CBMS) Continue engineering support.						
53) SSI NBCRS <i>FY 2010 Plans:</i> (CBMS) Initiate chemical biological capability sensor development using competitive prototyping. <i>FY 2011 Base Plans:</i> (CBMS) Continue chemical biological capability sensor development using competitive prototyping.		0.000	8.000	4.200	0.000	4.200
54) SSI NBCRS <i>FY 2010 Plans:</i> (CBMS) Initiate Chemical Biological (CB) capability sensor Developmental Test and Evaluation (DT&E) planning. <i>FY 2011 Base Plans:</i> (CBMS) Continue CB capability sensor DT&E planning and initiate T&E efforts.		0.000	0.224	0.955	0.000	0.955
55) SSI NBCRS <i>FY 2010 Plans:</i> (JCSD) Initiate engineering support, leveraging efforts performed under JNBCRS INC 3.		0.000	1.000	1.107	0.000	1.107

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2011 Base Plans:</i> (JCSD) Continue engineering support.						
56) SSI NBCRS <i>FY 2011 Base Plans:</i> (JCSD) Initiate sensor system development and demonstration using competitive prototyping.		0.000	0.000	8.080	0.000	8.080
57) SSI NBCRS <i>FY 2011 Base Plans:</i> (JCSD) Initiate sensor DT&E planning and initiate T&E efforts.		0.000	0.000	1.086	0.000	1.086
58) SSI NBCRS <i>FY 2011 Base Plans:</i> (SSI NBCRS) Initiate platform integration and system support of improved CB capable sensors for competitive prototype evaluation.		0.000	0.000	2.000	0.000	2.000
59) SSI NBCRS <i>FY 2010 Plans:</i> (SSI NBCRS) Initiate JPEO Integrated Program Assistance Team (IPAT) support for Stryker NBCRV path forward. <i>FY 2011 Base Plans:</i> (SSI NBCRS) Continue JPEO IPAT support for Stryker NBCRV path forward.		0.000	1.500	1.500	0.000	1.500
60) SSI NBCRS <i>FY 2010 Plans:</i> Congressional add for development of Man Portable Sensors for Dismounted Reconnaissance.		0.000	1.992	0.000	0.000	0.000

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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
61) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.	0.000	0.952	0.000	0.000	0.000
Accomplishments/Planned Programs Subtotals	46.316	78.042	124.936	0.000	124.936

C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011 Base</u>	<u>FY 2011 OCO</u>	<u>FY 2011 Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To Complete</u>	<u>Total Cost</u>
• JC0100: <i>JOINT BIO POINT DETECTION SYSTEM (JBPDS)</i>	75.545	41.976	43.555		43.555	41.252	52.776	73.164	71.894	Continuing	Continuing
• JC0101: <i>JS CHEM/BIO/RAD AGENT WATER MONITOR (JCBRAWM)</i>	6.000	3.184	0.000		0.000	0.000	0.000	0.000	0.000	Continuing	Continuing
• JC0250: <i>JOINT BIO STANDOFF DETECTOR SYSTEM (JBSDS)</i>	4.000	0.000	0.000		0.000	0.273	19.840	20.834	35.728	Continuing	Continuing
• JC1500: <i>NBC RECON VEHICLE (NBCRV)</i>	0.000	0.000	0.000		0.000	0.000	0.000	0.000	0.000	Continuing	Continuing
• JC4500: <i>NEXT GENERATION CHEMICAL STANDOFF DETECTION (NGCSD)</i>	0.000	0.000	0.000		0.000	0.000	9.840	12.120	21.799	Continuing	Continuing
• JF0100: <i>JOINT CHEMICAL AGENT DETECTOR (JCAD)</i>	58.406	27.694	40.071		40.071	45.805	52.762	53.330	63.217	Continuing	Continuing
• JN0900: <i>NON TRADITIONAL AGENT DETECTION (NTAD)</i>	0.000	0.000	4.178		4.178	4.075	3.376	6.745	8.918	Continuing	Continuing
• MC0100: <i>JOINT NBC RECONNAISSANCE SYSTEM (JNBCRS)</i>	32.699	32.421	22.511		22.511	65.779	122.214	50.385	0.000	Continuing	Continuing

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C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011 Base</u>	<u>FY 2011 OCO</u>	<u>FY 2011 Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To Complete</u>	<u>Total Cost</u>
• MC0101: <i>CBRN DISMOUNTED RECONNAISSANCE SYSTEMS (CBRN DRS)</i>	0.000	11.415	15.414		15.414	24.056	33.504	56.718	53.938	Continuing	Continuing
• MC0102: <i>JOINT CONTAMINATED SURFACE DETECTOR (JCSD)</i>	0.000	0.000	0.000		0.000	0.000	0.000	5.288	40.072	Continuing	Continuing
• MX0001: <i>JOINT BIO TACTICAL DETECTION SYSTEM (JBTD)</i>	0.000	0.000	0.000		0.000	8.080	19.060	29.237	33.933	Continuing	Continuing

D. Acquisition Strategy

CBRN DRS

The Chemical Biological Radiological Nuclear Dismounted Reconnaissance Systems (CBRN DRS) program uses spiral development with an evolutionary component/suite upgrade acquisition approach. Funding finalizes the design to provide the Services with enhanced full spectrum CBRN detection capability to support strategic, operational, and tactical objectives at lower life cycle costs. CBRN DRS will enhance the Situational Awareness (SA) by providing a dismounted ability to detect chemical, biological and radiological hazards across the Range of Military Operations (ROMO) and employ contamination avoidance activities to prevent disruption to operations and organizations.

JBPDS

The Joint Biological Point Detection System (JBPDS) uses an open systems approach to insert maturing and validated technologies as part of the overall acquisition strategy to expedite fielding of a credible force protection. The JBPDS Build 2 program will use results from a business case analysis to upgrade the system's line replaceable units (LRUs) to reduce life cycle costs, improve reliability, and address system obsolescence concerns. Per Director, Operational Test and Evaluation (DOT&E) Memorandum dated July 9, 2002, the program will continue to support the development of a Whole System Live Agent Test (WSLAT) capability.

JBSDS

The Joint Bio Stand-off Detector System (JBSDS) is employing an incremental acquisition strategy. JBSDS Increment 1 is the first standoff early warning biological detection (BD) system for the Joint Services. The JBSDS Increment 2 system will focus on providing 24-hour operations (Increment 1 is night-time only), improving

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<p>the false alarm rate and detection sensitivity, while decreasing size, weight and power. The JBSDS Increment 2 will also integrate with the global information network to provide near real time detection and warning theater-wide to limit the effect of biological agent hazards against U.S. forces at the tactical and operational levels of war. JBSDS Increment 3 will build on Increment 2 and focus on the development of a system that will operate on-the-move on mobile platforms as determined by the Warfighter.</p> <p>JBTDS</p> <p>The Joint Biological Tactical Detection (JBTDS) program will pursue an evolutionary incremental approach to provide capability to the Warfighter. The JBTDS program will develop, integrate, test, procure and field systems that improve biological aerosol detection, identification, and sampling capabilities. The JBTDS program will also reduce size, weight, power consumption, and logistic footprint over current systems. Test Readiness Evaluations (TRE) and Competitive Prototyping will support the JBTDS Engineering and Manufacturing Development (EMD) phase by identifying mature technologies and reducing overall risk. Modeling and simulation tools will be used in order to lower program risks, reduce costs and ensure a higher confidence in selected technologies.</p> <p>JCAD</p> <p>The current strategy employs a product improvement of the M4 JCAD to reduce Life Cycle costs, transition to a competitive procurement contract, and attain objective capability. Three competitive fixed-price contracts for the M4E1 were awarded in Sep 2007 for prototypes and options for full rate production. Competitive prototype testing was conducted and one system was selected for continued development. The production options will be exercised in FY11 following a successful production cut-in decision.</p> <p>JNBCRS 2</p> <p>The CBRN DRS (formerly JNBCRS Inc 2) fills a mission critical need to enhance Chemical, Biological, Radiological and Nuclear (CBRN) dismounted reconnaissance platoon capabilities. The program consists of two Phases. Phase I is the dismounted reconnaissance (DR) sets kits and outfits (SKO) configuration which provides an immediate critical need consisting of COTS and GOTS integrated into a modular, transportable container for dismounted operations. It will form the basis for Phase II which is the Monitoring and Survey (MS) SKO. The MS SKO will feature technology insertion, the addition of net-centric capability, and tailoring to focus on the service-specific needs, to include NTA detection.</p> <p>JNBCRS 3</p> <p>The JNBCRS Increment 3 program will develop and test system improvements to increase the military utility of the Stryker Product Improvement Program and Future Mounted Armored Reconnaissance Platforms. Separate Full and Open contracts will be awarded for both the CBMS Chem/Bio sensor and JCSD capabilities.</p>		

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<p>Competitively awarding these contracts will reduce the acquisition life cycle costs, weight, power requirements, and size for the reconnaissance platforms. The JCSD program transitioned from the CBRN Unmanned Ground Reconnaissance (CUGR) Advanced Concept Technology Demonstration (ACTD) into the Technology Development phase in FY09. The JNBCRS Increment 3 program supports transition to SSI NBCRS starting in FY10.</p> <p>JSLSCAD</p> <p>The acquisition strategy for the Joint Service Lightweight Standoff Chemical Agent Detector (JSLSCAD) production phase focused upon a dual path to procure required systems and concurrently develop and test system improvements to increase the military utility. The Milestone Decision Authority (MDA) approved procurement of additional JSLSCAD LRIP systems in February 2008. The Government awarded a Fixed Price Incentive contract to GD-ATP in July 2008 for production of systems to fulfill the Nuclear Biological Chemical Reconnaissance Vehicle (NBCRV) Extended LRIP requirements and additional delivery orders will be exercised for full rate production of systems to fulfill the remaining NBCRV requirements. The JSLSCAD program office awarded multiple contracts to support system engineering, software development, test and evaluation, and system support efforts to increase standoff detection capabilities to rapidly respond to evolving system integration requirements with minimal contractual lead time. All these efforts are being integrated into the Next Generation Chemical Standoff Detection (NGCSD) program.</p> <p>MDAP SPRT</p> <p>Major Defense Acquisition Program (MDAP) Support program will integrate System of Systems (SoS) solutions across the Armed Service's for Major Defense Acquisition Programs (MDAP) having Chemical and Biological Radiological and Nuclear (CBRN) survivability requirements. The MDAP program will achieve these SoS solutions by: (1) leading CBRN architecture development and System Engineering efforts that result in enterprise concepts that address requirements; (2) establishing agreements with the MDAPs on roles and responsibilities with respect to funding deliverables and integration; (3) demonstrating modular, net-centric, "plug and play" capabilities for mounted and dismounted CBRN reconnaissance requirements; (4) developing design and test schedules which synchronize support for CBRN capability integration with MDAPs' schedules; and (5) providing integrated program management across the CBRN commodity areas to deliver capabilities on time that support MDAP goals.</p> <p>NGCSD</p> <p>The Next Generation Chemical Standoff Detection (NGCSD) program, which was initiated under the JSLSCAD program, will award Indefinite Delivery/Indefinite Quantity contract(s) to support system engineering, software development, test and evaluation, and system support efforts to increase standoff detection capabilities. This contract type will allow the program office to rapidly respond to evolving system integration requirements and emerging test results with minimal contractual lead time. This will optimize the program goal of inserting the latest software and standoff detection technology into the host platforms in the shortest possible time.</p> <p>NTA DETECT</p>		

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<p>The Non-Traditional Agent (NTA) program will provide a detection capability through spiral evolutionary that will afford the Warfighter ability to attain situational awareness and respond to unknown and emerging hazards. The program provide a near term capability to detect priority emerging threat materials with common core technologies to detect and identify threats that can further be explored for lab deployable, fixed site and handheld applications. Leveraging COTS/GOTS assessments will be used in order to lower program risks, reduce costs, and ensure a higher confidence in selected technologies. The program will continue to address next priority mission areas and threats by continuing to qualify identified detection equipment.</p> <p>SSI NBCRS</p> <p>The Sensor Suite and Integration for Nuclear Biological Chemical Reconnaissance System (SSI NBCRS) program, transitioned from Joint Nuclear Biological Chemical Reconnaissance System (JNBCRS) Increment 3 in FY10, will develop and test platform specific prototype Chemical Biological Mass Spectrometer (CBMS) and Joint Contaminated Surface Detector (JCSD) capabilities. System development will be performed by separate full and open contract solicitations for CBMS and JCSD respectively, and will demonstrate a technology readiness level (TRL) of seven in laboratory and field testing. The contract efforts will finalize the technical approach and produce at least three prototypes of each system. Extensive laboratory and early user testing will be conducted prior to integration, test and evaluation into the JNBCRS. Upon successful completion of the JNBCRS integration, test and evaluation, a Milestone C In-Process Review (IPR) will be held to approve low-rate initial production of the CBMS and JCSD. The CBMS and JCSD will be introduced to the Stryker Fleets via Sensor Suite Improvements starting in FY15.</p> <p><u>E. Performance Metrics</u> N/A</p>		

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Product Development (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** CBRN DRS - DR SKO Program Development	C/CPIF	ICX Pittsburgh, PA	0.000	1.000	Jan 2010	4.500	Jan 2011	0.000		4.500	0.000	5.500	0.000
DR SKO Program Development	C/CPIF	ICX Pittsburgh, PA	0.000	1.400	Jan 2010	2.100	Jan 2011	0.000		2.100	0.000	3.500	0.000
DR SKO NTA Enhancements	C/FP	TBD	0.000	2.200	Jan 2010	15.023	Jul 2011	0.000		15.023	0.000	17.223	0.000
** CONG - Congressional Interest Item - Add Detection and Remediation of Bio/ Chem Weapons	C/CPFF	TBD	0.000	1.992	Jan 2010	0.000		0.000		0.000	0.000	1.992	0.000
** JBPDS - HW SB - Identifier development, modification and integration	C/CPFF	General Dynamics-Armament and Technical Charlotte, NC	0.000	2.740	Apr 2010	1.460	Jan 2011	0.000		1.460	0.000	4.200	0.000
HW SB - JBPDS Build II development, modification and integration	C/CPFF	General Dynamics-Armament and Technical Charlotte, NC	0.000	2.577	Apr 2010	0.905	Jan 2011	0.000		0.905	0.000	3.482	0.000
HW C - Development of new consumables for new Identifier	MIPR	JPM CBMS Ft. Detrick, MD	0.000	0.911	Oct 2009	1.229	Oct 2010	0.000		1.229	0.000	2.140	0.000
HW S - Hardware Cost - Purchase Build II Hardware for test	C/FFP	General Dynamics -	0.000	0.000		3.923	Jan 2011	0.000		3.923	0.000	3.923	0.000

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Product Development (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
		Armament and Technical Charlotte, NC											
HW SB - New Detector (RAAD) development, modification and integration	C/CPFF	General Dynamics - Armament & Technical Charlotte, NC	0.000	1.541	Jan 2010	1.389	Jan 2011	0.000		1.389	0.000	2.930	0.000
** MDAP SPRT - SW S - Develop Modular CBRN Sensing Capability	C/CPAF	TBD	0.000	1.528	Jan 2010	0.000		0.000		0.000	0.000	1.528	0.000
SW S - Decision Support Software Modeling and Simulation and Trade-Off Analysis	C/CPAF	TBD	0.000	1.683	Jan 2010	0.000		0.000		0.000	0.000	1.683	0.000
HW S - JSF Decon Shelter	MIPR	Various	0.000	1.800	Jan 2010	2.200	Jan 2011	0.000		2.200	0.000	4.000	0.000
HW S - JSF Decon	MIPR	Various	0.000	1.000	Jan 2010	3.805	Jan 2011	0.000		3.805	0.000	4.805	0.000
HW S - CBRN Sensor for SUGV	MIPR	Various	0.000	0.000		1.100	Jan 2011	0.000		1.100	0.000	1.100	0.000
** NGCSD - SW SB - Prototype System Development & Integration	C/CPFF	TBD	0.000	0.000		9.363	Jan 2011	0.000		9.363	0.000	9.363	0.000
SW C - System Integration Contract	C/CPFF	TBD	0.000	1.500	Jan 2010	0.000		0.000		0.000	0.000	1.500	0.000

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Product Development (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** NTA DETECT - HW S - DESI Mass Spec	C/CPAF	ICX Arlington, VA	0.000	2.475	Jan 2010	1.474	Jan 2011	0.000		1.474	0.000	3.949	0.000
HW S - GOTS/COTS Dual Use Assessment	C/CPAF	Battelle Crystal City, VA	0.000	1.104	Jan 2010	1.000	Jan 2011	0.000		1.000	0.000	2.104	0.000
SW S - DESI Mass Spec Library Development	MIPR	RDECOM Aberdeen Proving Ground, MD	0.000	0.950	Oct 2009	0.370	Oct 2010	0.000		0.370	0.000	1.320	0.000
HW S - COTS Enzyme based technologies	C/CPAF	Agentase - ICX Pittsburgh, PA	0.000	0.000		1.100	Jan 2011	0.000		1.100	0.000	1.100	0.000
** SSI NBCRS - SW SB - (CBMS) Chemical Biological Sensor Capability Development	C/CPIF	TBD	0.000	8.000	Apr 2010	4.200	Jan 2011	0.000		4.200	0.000	12.200	0.000
HW C - (JSCD) Sensor System Development and Demonstration	C/CPIF	TBD	0.000	0.000		8.080	Jan 2011	0.000		8.080	0.000	8.080	0.000
HW S - (SSI NBCRS) Sensor Platform Integration	C/CPIF	TBD	0.000	0.000		2.000	Jan 2011	0.000		2.000	0.000	2.000	0.000
HW C - Develop Man Portable Sensors for Dismounted Reconnaissance	C/CPIF	TBD	0.000	1.992	Jan 2010	0.000		0.000		0.000	0.000	1.992	0.000
Subtotal			0.000	36.393		65.221		0.000		65.221	0.000	101.614	0.000

Remarks

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Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** CBRN DRS - ES S - NTA Enhancements	C/CPIF	Various TBD	0.000	0.000		3.860	Jan 2011	0.000		3.860	0.000	3.860	0.000
** JBPDS - ILS SB - New Identifier/Collector/ Detector logistics and support documentation	C/CPFF	General Dynamics - Armament and Technical Charlotte, NC	0.000	0.671	Jan 2010	2.213	Jan 2011	0.000		2.213	0.000	2.884	0.000
** NGCSD - TD/D SB - Logistics Planning and Development	MIPR	Various	0.000	0.000		0.600	Jan 2011	0.000		0.600	0.000	0.600	0.000
** NTA DETECT - ES SB - Mass Spectrometer Analysis and Evaluation	PO	TBD	0.000	0.675	Oct 2009	0.325	Oct 2010	0.000		0.325	0.000	1.000	0.000
Subtotal			0.000	1.346		6.998		0.000		6.998	0.000	8.344	0.000

Remarks

Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** CBRN DRS - DTE S - DR SKO	MIPR	ATEC Alexandria, VA	0.000	2.300	Jan 2010	2.696	Jan 2011	0.000		2.696	0.000	4.996	0.000

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Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
DTE S - DR SKO Developmental Testing	MIPR	ATEC Alexandria, VA	0.000	2.819	Jan 2010	2.800	Jan 2011	0.000		2.800	0.000	5.619	0.000
DTE S - NTA Enhancements	MIPR	ATEC Alexandria, VA	0.000	0.000		11.800	Jan 2011	0.000		11.800	0.000	11.800	0.000
** JBPDS - DTE SB - New Identifier/Collector/ Detector developmental testing	C/CPFF	General Dynamics - Armament and Technical Charlotte, NC	0.000	1.689	Jan 2010	3.513	Jan 2011	0.000		3.513	0.000	5.202	0.000
OTE C - Identifier consumable testing	MIPR	JPM CBMS Ft Detrick, MD	0.000	0.450	Oct 2009	0.950	Oct 2010	0.000		0.950	0.000	1.400	0.000
** JCAD - DTE S - M4E1 JCAD Developmental Test	MIPR	Various	34.815	3.360	Jan 2010	0.000		0.000		0.000	0.000	38.175	0.000
OTE S - M4E1 JCAD Operational Test and Evaluation	MIPR	Various	4.980	1.510	Jan 2010	0.000		0.000		0.000	0.000	6.490	0.000
OTE S - M4E1 Multi-Service Operational Test and Evaluation	MIPR	Various	0.000	0.000		8.000	Jan 2011	0.000		8.000	0.000	8.000	0.000
** NGCSD - OTHT S - Test Methodology Development	MIPR	Various	0.000	3.615	Apr 2010	0.000		0.000		0.000	0.000	3.615	0.000
** NTA DETECT - DTE S - Developmental Test Mass Spectrometer	MIPR	ECBC APG, MD	0.000	8.252	Oct 2009	2.400	Oct 2010	0.000		2.400	0.000	10.652	0.000

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Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** SSI NBCRS - OTHT SB - (CBMS) Developmental Testing	MIPR	Various	0.000	0.224	Jan 2010	0.955	Jan 2011	0.000		0.955	0.000	1.179	0.000
DTE C - (JCSD) Developmental Testing	MIPR	Various	0.000	0.000		1.086	Jan 2011	0.000		1.086	0.000	1.086	0.000
Subtotal			39.795	24.219		34.200		0.000		34.200	0.000	98.214	0.000

Remarks

Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** CBRN DRS - PM/MS-S - Program Management and System Engineering Support	PO	JPM NBC CA APG, MD	0.000	0.718	Oct 2009	1.407	Oct 2010	0.000		1.407	0.000	2.125	0.000
PM/MS S - NTA Enhancements Program Management and System Engineering Support	PO	JPM NBC CA APG, MD	0.000	0.000		1.200	Oct 2010	0.000		1.200	0.000	1.200	0.000
	MIPR	JPM BD	2.446	0.938	Oct 2009	0.886	Oct 2010	0.000		0.886	0.000	4.270	0.000

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Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** JBPDS - PM/MS S - Project Management		APG, MD											
PM/MS S - Project Management, JPEO MGMT	PO	JPEO CBD Falls Church, VA	0.000	0.982	Jan 2010	0.914	Jan 2011	0.000		0.914	0.000	1.896	0.000
** JCAD - PM/MS S - Joint Service Support	MIPR	Various	2.998	3.206	Jan 2010	1.733	Jan 2011	0.000		1.733	0.000	7.937	0.000
** MDAP SPRT - PM/MS SB - Management & Oversight	MIPR	Various	0.000	3.200	Jan 2010	2.155	Jan 2011	0.000		2.155	0.000	5.355	0.000
** NGCSD - PM/MS S - Program Management and Systems Engineering Support	MIPR	JPM NBC CA APG, MD	0.000	0.000		1.500	Oct 2010	0.000		1.500	0.000	1.500	0.000
PM/MS SB - Joint Service Combat Developer Support	MIPR	Various	0.000	0.000		0.400	Jan 2011	0.000		0.400	0.000	0.400	0.000
PM/MS S - Program Management and Systems Engineering Support	MIPR	JPEO-CBD Falls Church, VA	0.000	1.500	Jan 2010	0.750	Oct 2009	0.000		0.750	0.000	2.250	0.000
** NTA DETECT - PM/MS S - Program Management support	PO	JPEO Falls Church, VA	0.000	0.904	Jul 2010	3.857	Jul 2011	0.000		3.857	0.000	4.761	0.000
** SSI NBCRS - PM/MS S - (CBMS) Program	MIPR	JPM NBC CA APG, MD	0.000	1.184	Oct 2009	1.108	Oct 2010	0.000		1.108	0.000	2.292	0.000

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Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
Management and Systems Engineering Support													
PM/MS S - (JCSD) Program Management and Systems Engineering Support	MIPR	JPM NBC CA APG, MD	0.000	1.000	Oct 2009	1.107	Oct 2010	0.000		1.107	0.000	2.107	0.000
PM/MS S - (SSI NBCRS) Program Management and Systems Engineering Support	MIPR	JPEO-CBD Falls Church, VA	0.000	1.500	Jan 2010	1.500	Oct 2010	0.000		1.500	0.000	3.000	0.000
** ZSBIR - SBIR/STTR - Aggregated from ZSBIR-SBIR/STTR	MIPR	HQ AMC, Alexandria	0.000	0.952		0.000		0.000		0.000	0.000	0.952	0.000
Subtotal			5.444	16.084		18.517		0.000		18.517	0.000	40.045	0.000

Remarks

	Total Prior Years Cost	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
Project Cost Totals	45.239	78.042	124.936	0.000	124.936	0.000	248.217	0.000

Remarks

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT CA5: <i>CONTAMINATION AVOIDANCE (SDD)</i>
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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
** CBRN DRS - Dismounted Reconnaissance (DR) Preliminary Design Review		■																										
Dismounted Reconnaissance (DR) Prototype Development and Test		■	■	■	■	■	■																					
Dismounted Reconnaissance (DR) Milestone (MS) C LRIP								■																				
Monitoring and Survey (MS) CPD																	■											
Monitoring and Survey (MS) Milestone C																			■									
Monitoring and Survey (MS) LRIP																					■	■	■	■	■	■	■	
** CONG - Self contained automated vehicle washing systems with microwave decontamination								■	■	■	■	■	■															
Protective Self-Decontaminating Surfaces - CHRPS								■	■	■	■	■	■															
** JBPDS - MS C Full Rate Production Decision (FRP)				■																								
FRP Contract Award								■																				
Full Rate Production (First Full Contract Award)								■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	
Build II - Development and Integration					■	■	■	■	■	■	■	■	■	■	■	■	■											
Build II - Test and validation of LRU improvements																	■	■	■	■								
Whole System Live Agent Test II																	■	■	■	■								

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
** JBSDS - Increment I JBSDS LRIP 2	■	■	■	■	■	■																						
Increment 1 JBSDS Full Material Release						■																						
Increment 1 JBSDS First Unit Equipped (FUE)						■																						
Increment 1 - JBSDS FRP						■																						
Increment 2 - Technology Modeling	■	■	■	■	■	■	■																					
Increment 2 - Pre-Milestone A	■	■	■	■	■	■	■																					
Increment 2 - Milestone A							■																					
Increment 2 - Technology Development							■	■	■	■	■	■	■	■														
Increment 2 - Preliminary Design Review													■															
Increment 2 - Milestone B															■													
Increment 2 - Engineering & Manufacturing Development															■	■	■	■	■	■	■	■	■	■	■	■		
Increment 2 - Milestone C																										■		
Increment 2 - LRIP																										■	■	■
** JBTDS - Materiel Development Decision				■																								
MS A Decision							■																					
Competitive Prototyping Contract Award								■																				
Competitive Prototyping Testing										■	■																	
Capability Development Document										■																		
PDR											■																	

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
MS B Decision												■																
EMD Contract Award												■																
Developmental Testing												■	■	■	■	■	■	■	■									
CDR															■													
MS C Decision																												
MS C Contract Award																												
Production Verification Test																												
IOT&E																												
FRP Decision																												
FRP Contract Award #2																												
IOC																												
** JCAD - M4E1 JCAD - Customer Testing				■	■																							
M4E1 JCAD - Developmental Testing						■	■	■																				
M4E1 JCAD - Operational Testing								■																				
M4E1 JCAD - Production Cut-in Decision												■																
** JNBCRS 2 - JNBCRS INC 2 - DR SKO Milestone C Low Rate Initial Production								■	■	■	■	■	■	■	■													
** JNBCRS 3 - JNBCRS INC 3 (JCSD) - Hardware Maturation Effort	■	■																										
JNBCRS INC 3 - Development Testing		■	■	■	■																							
JNBCRS INC 3 - Sensor Development				■	■	■	■																					

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
** JSLSCAD - SoS Technology Demo	■																											
SoS Program	■	■	■	■																								
SoS Operational Demo			■	■																								
** MDAP SPRT - System of Systems (SoS) Component Development	■	■	■	■	■	■	■	■	■	■	■	■																
Data Fusion Algorithm Development	■	■	■	■	■	■	■	■																				
Collective Protection Advanced Technology Demonstrator Developmental Test (DT)							■	■																				
Reactive/Removable Coating Developmental Test (DT)						■	■	■																				
Catox Tech Demonstration for Abrams Main Battle Tank					■	■	■	■	■	■	■	■																
Advance Component Prototype Development of JSF Decontamination			■	■	■	■	■	■	■	■	■	■																
Modular Individual Protection Design and Test									■	■	■	■	■	■														
** NGCSD - Material Development Decision (MDD)						■																						
Analysis of Alternatives (AoA)							■	■																				
MS A										■																		
Competitive Prototyping										■	■	■	■															
Preliminary Design Review (PDR)													■															
MS B																											■	

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program			DATE: February 2010
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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
DT													■	■	■	■	■	■	■	■								
Milestone C - LRIP																					■							
LRIP																					■	■	■	■				
Operational Testing																												■
** NTA DETECT - COTS/GOTS DT/MUA					■	■	■																					
COTS/GOTS Interim Capability							■	■	■																			
Lab Deployable Mass Spec DT/OA					■	■	■	■																				
Lab Deployable Mass Spec Transition												■																
Man Portable Mass Spec DT/OA										■	■	■	■															
Man Portable Mass Spec Transition													■															
Man Portable Mass Spec Integration																■												
Aerosol Detection DT										■	■	■																
Aerosol Detection OA																■												
** SSI NBCRS - Prototype Sensor Technology Evaluation							■	■	■	■																		
Prototype Sensor Developmental Testing and Evaluation										■	■	■	■															
(JCS) PDR IPR																■												
(CBMS) PDR IPR																■												
Engineering & Manufacturing Development (EMD) Sensor Platform Integration																	■	■	■	■	■	■	■	■				

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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015							
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4				
Platform Developmental Testing and Evaluation																					■	■	■									
(CBMS & JCSD) LRIP IPR																															■	

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
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Schedule Details

Event	Start		End	
	Quarter	Year	Quarter	Year
** CBRN DRS - Dismounted Reconnaissance (DR) Preliminary Design Review	2	2009	2	2009
Dismounted Reconnaissance (DR) Prototype Development and Test	2	2009	3	2010
Dismounted Reconnaissance (DR) Milestone (MS) C LRIP	4	2010	4	2010
Monitoring and Survey (MS) CPD	4	2013	4	2013
Monitoring and Survey (MS) Milestone C	3	2014	3	2014
Monitoring and Survey (MS) LRIP	4	2014	2	2016
** CONG - Self contained automated vehicle washing systems with microwave decontamination	3	2010	4	2011
Protective Self-Decontaminating Surfaces - CHRPS	3	2010	4	2011
** JBPDS - MS C Full Rate Production Decision (FRP)	4	2009	4	2009
FRP Contract Award	3	2010	3	2010
Full Rate Production (First Full Contract Award)	3	2010	4	2016
Build II - Development and Integration	1	2010	3	2013
Build II - Test and validation of LRU improvements	1	2013	4	2013
Whole System Live Agent Test II	1	2013	4	2013
** JBSDS - Increment I JBSDS LRIP 2	2	2008	2	2010
Increment 1 JBSDS Full Material Release	2	2010	2	2010
Increment 1 JBSDS First Unit Equipped (FUE)	2	2010	2	2010
Increment 1 - JBSDS FRP	2	2010	2	2010

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Event	Start		End	
	Quarter	Year	Quarter	Year
Increment 2 - Technology Modeling	4	2004	3	2010
Increment 2 - Pre-Milestone A	1	2008	3	2010
Increment 2 - Milestone A	3	2010	3	2010
Increment 2 - Technology Development	3	2010	3	2012
Increment 2 - Preliminary Design Review	2	2012	2	2012
Increment 2 - Milestone B	3	2012	3	2012
Increment 2 - Engineering & Manufacturing Development	3	2012	1	2015
Increment 2 - Milestone C	1	2015	1	2015
Increment 2 - LRIP	1	2015	2	2017
** JBTDs - Materiel Development Decision	4	2009	4	2009
MS A Decision	3	2010	3	2010
Competitive Prototyping Contract Award	4	2010	4	2010
Competitive Prototyping Testing	1	2011	2	2011
Capability Development Document	1	2011	1	2011
PDR	2	2011	2	2011
MS B Decision	3	2011	3	2011
EMD Contract Award	3	2011	3	2011
Developmental Testing	3	2011	3	2013
CDR	4	2012	4	2012
MS C Decision	3	2013	3	2013

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Event	Start		End	
	Quarter	Year	Quarter	Year
MS C Contract Award	3	2013	3	2013
Production Verification Test	1	2014	1	2014
IOT&E	1	2014	3	2014
FRP Decision	3	2014	3	2014
FRP Contract Award #2	3	2014	3	2014
IOC	3	2015	3	2015
** JCAD - M4E1 JCAD - Customer Testing	4	2009	1	2010
M4E1 JCAD - Developmental Testing	2	2010	4	2010
M4E1 JCAD - Operational Testing	4	2010	4	2010
M4E1 JCAD - Production Cut-in Decision	2	2011	2	2011
** JNBCRS 2 - JNBCRS INC 2 - DR SKO Milestone C Low Rate Initial Production	3	2010	2	2012
** JNBCRS 3 - JNBCRS INC 3 (JCSD) - Hardware Maturation Effort	1	2008	2	2009
JNBCRS INC 3 - Development Testing	2	2009	1	2010
JNBCRS INC 3 - Sensor Development	4	2009	3	2010
** JSLSCAD - SoS Technology Demo	4	2008	1	2009
SoS Program	4	2008	4	2009
SoS Operational Demo	3	2009	4	2009
** MDAP SPRT - System of Systems (SoS) Component Development	2	2007	4	2011
Data Fusion Algorithm Development	2	2007	4	2010
Collective Protection Advanced Technology Demonstrator Developmental Test (DT)	3	2010	4	2010

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Event	Start		End	
	Quarter	Year	Quarter	Year
Reactive/Removable Coating Developmental Test (DT)	2	2010	4	2010
Catox Tech Demonstration for Abrams Main Battle Tank	1	2010	4	2011
Advance Component Prototype Development of JSF Decontamination	4	2009	1	2012
Modular Individual Protection Design and Test	1	2011	2	2012
** NGCSD - Material Development Decision (MDD)	2	2010	2	2010
Analysis of Alternatives (AoA)	3	2010	4	2010
MS A	1	2011	1	2011
Competitive Prototyping	1	2011	4	2011
Preliminary Design Review (PDR)	4	2011	4	2011
MS B	1	2012	1	2012
DT	2	2012	4	2013
Milestone C - LRIP	1	2014	1	2014
LRIP	1	2014	4	2014
Operational Testing	1	2015	1	2015
** NTA DETECT - COTS/GOTS DT/MUA	1	2010	3	2010
COTS/GOTS Interim Capability	3	2010	1	2011
Lab Deployable Mass Spec DT/OA	1	2010	4	2010
Lab Deployable Mass Spec Transition	4	2011	4	2011
Man Portable Mass Spec DT/OA	3	2011	2	2012
Man Portable Mass Spec Transition	2	2012	2	2012

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Event	Start		End	
	Quarter	Year	Quarter	Year
Man Portable Mass Spec Integration	3	2013	3	2013
Aerosol Detection DT	3	2011	1	2012
Aerosol Detection OA	1	2013	1	2013
** SSI NBCRS - Prototype Sensor Technology Evaluation	3	2010	2	2011
Prototype Sensor Developmental Testing and Evaluation	3	2011	2	2012
(JCSD) PDR IPR	4	2012	4	2012
(CBMS) PDR IPR	4	2012	4	2012
Engineering & Manufacturing Development (EMD) Sensor Platform Integration	1	2013	4	2014
Platform Developmental Testing and Evaluation	1	2014	3	2014
(CBMS & JCSD) LRIP IPR	2	2015	2	2015

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program									DATE: February 2010		
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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
CM5: <i>HOMELAND DEFENSE (SDD)</i>	3.897	8.638	1.166	0.000	1.166	3.822	0.000	2.361	2.413	Continuing	Continuing
Quantity of RDT&E Articles	0	0	3		3	0	0	0	0		

A. Mission Description and Budget Item Justification

The PM Consequence Management program supports the development of a Common Analytical Laboratory System capability (CALs) that will be modular, scalable and adaptable to a variety of CONOPS and environmental conditions. Currently, fielded systems have been designed independently by various agencies with the intent of meeting a specific units requirements. As a result, multiple mobile lab configurations exist with differing sustainment tails and lacking in commonality. The system under development will incorporate an open architecture that can accommodate quick installation or removal of equipment as mission requirements dictate. As well, it will provide the ability to rapidly develop a common operating picture allowing first responders and DoD officials to determine the appropriate course of action. The analytical detection package fielded will be fitted to the specific mission and CONOPS of the gaining unit and be able to detect and identify Chemical Warfare Agents (CWAs), Toxic Industrial Chemicals (TICs), Toxic Industrial Materials (TIMs), Biological Warfare Agents (BWAs), Lower Explosive Limits (LEL), and radioactive particles in all sample types.

The CB Installation Protection Program (CBIPP) supports the development of analytical methodologies to expand/enhance the operational capabilities of currently fielded CBRN detection, identification and protection technologies against emerging threats to include Toxic Industrial Chemicals (TICs), Chemical Warfare Agents (CWAs), and Biological Warfare Agents (BWAs). Detection and identification of these substances is currently difficult and time-consuming. Current systems lack extensive libraries to support rapid identification. Identification may also involve multiple, expensive technologies. The ability to rapidly detect and identify a TIC is essential to effectively control and mitigate its effects, thus protecting personnel. This program also supports the evaluation of emerging CBRN detection, identification, information management and decision support technologies to DoD response units to maintain required state of the art capabilities.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) CALS	0.000	0.499	0.150	0.000	0.150

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010				
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B. Accomplishments/Planned Program (\$ in Millions)								
				FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2010 Plans:</i> Initiate Program Office planning and programming. <i>FY 2011 Base Plans:</i> Continue Program Office planning and programming.								
2) CALS <i>FY 2010 Plans:</i> Initiate System Engineering and Logistics Support. <i>FY 2011 Base Plans:</i> Continue System Engineering and Logistics Support.				0.000	0.971	0.150	0.000	0.150
3) CALS <i>FY 2010 Plans:</i> Initiate Subsystem Design and Development - Open Architecture Design Analytics and Laboratory Information Management. <i>FY 2011 Base Plans:</i> Complete Subsystem Design and Development - Open Architecture Design Analytics and Laboratory Information Management.				0.000	3.464	0.000	0.000	0.000
4) CALS <i>FY 2010 Plans:</i> Initiate/conduct Developmental Testing.				0.000	0.732	0.566	0.000	0.566

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2011 Base Plans:</i> Conduct Developmental Testing.						
5) CALS <i>FY 2011 Base Plans:</i> Conduct system prototype refurbishment and refinement.		0.000	0.000	0.300	0.000	0.300
6) CBIPP <i>FY 2009 Accomplishments:</i> Supported development of analytical methodologies to expand CBRN detection, identification, and protection capabilities.		0.750	0.000	0.000	0.000	0.000
7) CBIPP <i>FY 2010 Plans:</i> Supports the development of methodologies used to perform CBRN detection and evaluation under various environmental conditions.		0.000	1.091	0.000	0.000	0.000
8) CBIPP <i>FY 2010 Plans:</i> Supports the evaluation of CBRN detection, identification, information management and decision support technologies.		0.000	1.770	0.000	0.000	0.000
9) CBIPP <i>FY 2009 Accomplishments:</i> Supported the evaluation of CBRN detection, identification, information management, and decision support technologies.		1.697	0.000	0.000	0.000	0.000

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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
10) WMD CST <i>FY 2009 Accomplishments:</i> Conducted Program Office planning and programming.	0.450	0.000	0.000	0.000	0.000
11) WMD CST <i>FY 2009 Accomplishments:</i> Conducted Systems Engineering and Logistics Support.	1.000	0.000	0.000	0.000	0.000
12) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.	0.000	0.111	0.000	0.000	0.000
Accomplishments/Planned Programs Subtotals					
	3.897	8.638	1.166	0.000	1.166

C. Other Program Funding Summary (\$ in Millions)

Line Item	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	FY 2012	FY 2013	FY 2014	FY 2015	Cost To Complete	Total Cost
• JS0004: <i>WMD - CIVIL SUPPORT TEAMS (WMD CST)</i>	8.300	11.765	39.862		39.862	33.402	37.398	44.817	47.159	Continuing	Continuing
• JS0500: <i>CB INSTALLATION/ FORCE PROTECTION PROGRAM (FORCE PROT)</i>	80.103	53.623	50.773		50.773	60.324	59.836	57.840	54.455	Continuing	Continuing

D. Acquisition Strategy

CALS

The Common Analytical Laboratory System (CALs) will follow an incremental approach designed to address known joint force capability requirements for Chemical, Biological, Radiological and Nuclear (CBRN) detection which includes Toxic Industrial Chemicals (TICs), Toxic Industrial Materials (TIMs), Chemical Warfare Agents

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT CM5: <i>HOMELAND DEFENSE (SDD)</i>
<p>(CWAs), Biological Warfare Agents (BWAs). CALS will address situational awareness by leveraging efforts underway with JPEO-CBD to the extent possible. CALS will accommodate these component requirements within a modular and scalable concept framework.</p> <p>FORCE PROT</p> <p>The Special Study for System Methodology Development will support the development of analytical methodologies to expand/enhance the operational capabilities of currently fielded CBRN detection, identification and protection technologies against emerging threats to include TIC, CWA, and BWA threats.</p> <p>The Special Study for CBRN Defense Technology Evaluation will support the evaluation of emerging CBRN detection, identification, information management and decision support technologies to DoD response units to maintain required state-of-the-art capabilities.</p> <p>WMD CST</p> <p>This program utilizes multiple acquisition vehicles to deliver a CBRN capability to the WMD response units. The CALS program will upgrade the analytical capability with the objective of improving chemical and biological detection sensitivity and selectivity of the WMD CST Analytical Laboratory System Increment 1 and the 20th SUPCOM heavy and light tactical lab variants. Additionally, the CALS will integrate the communications and reachback capability for mobile CBRN homeland defense capability as required by the JROC.</p> <p><u>E. Performance Metrics</u> N/A</p>		

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT CM5: <i>HOMELAND DEFENSE (SDD)</i>
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Product Development (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** CALS - HW SB - Analytical Subsystem Design and Development	C/CPIF	TBD	0.000	2.252	Jan 2010	0.000		0.000		0.000	0.000	2.252	0.000
SW SB - Information Technology Subsystem Development	C/CPIF	TBD	0.000	1.042	Jan 2010	0.000		0.000		0.000	0.000	1.042	0.000
HW S - Analytical Protocol Development	MIPR	TBD	0.000	0.170	Apr 2010	0.000		0.000		0.000	0.000	0.170	0.000
HW S - CALS Prototype Systems	C/CPIF	TBD	0.000	0.000		0.300	Jul 2011	0.000		0.300	0.000	0.300	0.000
** FORCE PROT - HW S - System Protocol Development	C/FP	TBD	0.000	1.091	Oct 2009	0.000		0.000		0.000	0.000	1.091	0.000
Subtotal			0.000	4.555		0.300		0.000		0.300	0.000	4.855	0.000

Remarks

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** CALS - ES S - Engineering Support	MIPR	Edgewood Chemical and Biological Center Edgewood, MD	0.000	0.499	Oct 2009	0.150	Jan 2011	0.000		0.150	0.000	0.649	0.000
Subtotal			0.000	0.499		0.150		0.000		0.150	0.000	0.649	0.000

Remarks

Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** CALS - DTE SB - Analytical System Developmental Testing	MIPR	TBD	0.000	0.732	Apr 2010	0.000		0.000		0.000	0.000	0.732	0.000
DTE S - System Developmental Testing	MIPR	TBD	0.000	0.000		0.566	Jul 2011	0.000		0.566	0.000	0.566	0.000
** FORCE PROT - OTHT C - System Component Testing	C/FP	TBD	0.000	1.770	Oct 2009	0.000		0.000		0.000	0.000	1.770	0.000
Subtotal			0.000	2.502		0.566		0.000		0.566	0.000	3.068	0.000

Remarks

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** CALS - PM/MS HW - Program Office - Planning and Programming	MIPR	Edgewood Chemical Biological Center Edgewood, MD	0.000	0.971	Oct 2009	0.150	Jan 2011	0.000		0.150	0.000	1.121	0.000
** ZSBIR - SBIR/STTR - Aggregated from ZSBIR- SBIR/STTR	MIPR	HQ AMC, Alexandria	0.000	0.111		0.000		0.000		0.000	0.000	0.111	0.000
Subtotal			0.000	1.082		0.150		0.000		0.150	0.000	1.232	0.000

Remarks

	Total Prior Years Cost	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
Project Cost Totals	0.000	8.638	1.166	0.000	1.166	0.000	9.804	0.000

Remarks

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT CM5: <i>HOMELAND DEFENSE (SDD)</i>
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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
** CALS - CALS Program Initiation					■																							
CALS Design, Development and Integration					■	■	■	■	■	■	■	■	■	■														
CALS System Demonstration																■												
CALS MDD					■																							
CALS Milestone A								■																				
CALS Milestone C																■												
** FORCE PROT - System Methodologies Development	■	■	■	■																								
Technology Evaluation	■	■	■	■																								
System Architecture Development					■	■	■	■																				
Bio-Collection/Detection Evaluation					■	■	■	■																				
** WMD CST - CALS Program Initiation					■																							
CALS Design, Development and Integration #2					■	■	■	■	■	■	■	■	■	■														
CALS System Demonstration #2																■												
CALS MDD #2					■																							
CALS Milestone A #2					■																							

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT CM5: <i>HOMELAND DEFENSE (SDD)</i>

Schedule Details

Event	Start		End	
	Quarter	Year	Quarter	Year
** CALS - CALS Program Initiation	1	2010	1	2010
CALS Design, Development and Integration	1	2010	2	2012
CALS System Demonstration	2	2012	2	2012
CALS MDD	1	2010	1	2010
CALS Milestone A	3	2010	3	2010
CALS Milestone C	2	2012	2	2012
** FORCE PROT - System Methodologies Development	1	2009	4	2009
Technology Evaluation	1	2009	4	2009
System Architecture Development	1	2010	4	2010
Bio-Collection/Detection Evaluation	1	2010	4	2010
** WMD CST - CALS Program Initiation	1	2010	1	2010
CALS Design, Development and Integration #2	1	2010	2	2012
CALS System Demonstration #2	2	2012	2	2012
CALS MDD #2	1	2010	1	2010
CALS Milestone A #2	1	2010	1	2010

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT CO5: <i>COLLECTIVE PROTECTION (SDD)</i>
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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
CO5: <i>COLLECTIVE PROTECTION (SDD)</i>	13.323	12.002	18.459	0.000	18.459	11.671	10.267	7.835	0.000	Continuing	Continuing
Quantity of RDT&E Articles	0	72	0		0	0	72	0	0		

A. Mission Description and Budget Item Justification

Funding supports System Development and Demonstration and Low Rate Initial Production (SDD/LRIP) of Joint Service Chemical, Biological, and Radiological (CBR) Collective Protection (CP) systems that are smaller, lighter, less costly to produce and maintain, and more logistically supportable enabling mission accomplishment in CBR environments. CP systems can be installed on any type of platform, such as, hard and soft shelters, vehicles, ships, aircraft, and buildings. CP systems provide spaces safe from the effects of CBR contamination.

Systems funded under this project are: Joint Expeditionary Collective Protection (JECP).

JECP provides the Joint Expeditionary Forces a CP capability which is lightweight, compact, modular, and affordable. A family of systems is planned that will allow the application of CP to transportable soft-side shelters, enclosed spaces of opportunity, and in remote austere locations as a standalone resource. JECP will be capable of protecting personnel groups of varying size, unencumbered by Individual Protective Equipment (IPE), from the effects of CB agents, Toxic Industrial Materials (TIMs), radiological particles, heat, dust, and sand. The employment of JECP is a strategic deterrence against enemy use of CBR agents or TIMs, and will reduce the need for personnel and equipment decontamination.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) JECP - Engineering and Manufacturing Development (EMD) Contract Engineering and Manufacturing Development Contract to design, develop, integrate and test the prototype Joint Expeditionary Collective Protection (JECP) Family of Systems (FoS) that meet the requirements of the Capability Development Document (CDD) and System Performance Specification (SPS).	3.630	3.212	1.330	0.000	1.330

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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2009 Accomplishments:</i> Established the Functional baseline by developing and placing the System Performance Specification (SPS) and the Requirements Traceability Management (RTM) under configuration control. Planned and conducted the System Functional Review. Identified new risks and managed existing risks as delegated. Identified Environmental Safety Occupational Health (ESOH) hazards and managed corresponding ESOH hazard risks. Assisted in HSI assessments and reviews.</p> <p><i>FY 2010 Plans:</i> Update and maintain the RTM to track when requirements have been verified as test results become available. Participate in Configuration Control Board as the detailed design evolves and drawings are developed and updated.</p> <p><i>FY 2011 Base Plans:</i> Develop, update and/or review program documentation in preparation for MS C (as documented above for MS B).</p>					
<p>8) JECIP - Test and Evaluation IPT Lead and oversee all aspects of the JECIP Integrated Test (IT) program</p> <p><i>FY 2009 Accomplishments:</i> Identified requirements for a System Performance Model (SPM) through development of the JECIP SPM Software Requirement Spec. Coordinated model development. Conducted integrated test planning, coordination, and test readiness reviews associated with contractor and government component level testing. Developed and/or reviewed test plans, procedures and reports and participated in test events as needed. Developed and/or reviewed Design of Experiment and associated Test Matrices in support of government component level DT and models as necessary. Reviewed contractor component level test data and reports (including reliability test results and KPP failures).</p>	1.100	0.940	1.000	0.000	1.000

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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2010 Plans:</i> Conduct integrated test planning, coordination, and test readiness reviews associated with government component level testing and contractor and government system level testing. Develop test plans, procedures and reports and participate in test events as needed. Develop Design of Experiment and associated Test Matrices in support of government system level testing. Review component level test data and reports (including reliability test results and KPP failures).</p> <p><i>FY 2011 Base Plans:</i> Develop and/or review test plans, procedures and reports and participate in government system level DT and combined DT/OT events as needed. Develop, update and/or review program documentation in preparation for MS C (as documented above for MS B).</p>					
<p>9) JECF - Integrated Logistics Support IPT</p> <p>Oversee and provide supportability planning guidance to the EMD contractor in addressing logistic support elements including maintenance philosophy, manpower & personnel, supply support, Tech Data, support & test equipment, training and training support.</p> <p><i>FY 2009 Accomplishments:</i> Initiated a Business Case Analysis (BCA) to determine the scope of implementing Performance Based Logistics (PBL) and evaluate whether organic or Contractor Logistics Support is the most effective approach.</p> <p><i>FY 2010 Plans:</i> Continue the Business Case Analysis (BCA) to determine the scope of implementing PBL. Continue evaluation as to whether organic or Contractor Logistics Support is the most effective approach. Begin an analysis to identify surge requirements and industries ability to support.</p>	0.469	0.650	0.500	0.000	0.500

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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2010 Plans:</i> Provide strategic tactical planning, government systems engineering, program/financial management, costing, technology assessment, contracting, scheduling, acquisition oversight and technical support.</p> <p><i>FY 2011 Base Plans:</i> Provide strategic tactical planning, government systems engineering, program/financial management, costing, technology assessment, contracting, scheduling, acquisition oversight and technical support.</p>					
12) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.	0.000	0.155	0.000	0.000	0.000
Accomplishments/Planned Programs Subtotals					
	13.323	12.002	18.459	0.000	18.459

C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011 Base</u>	<u>FY 2011 OCO</u>	<u>FY 2011 Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To Complete</u>	<u>Total Cost</u>
• JN0014: <i>COLLECTIVE PROT SYS AMPHIB BACKFIT (CPS BKFT)</i>	18.219	11.963	5.869		5.869	0.000	0.000	0.000	0.000	Continuing	Continuing
• JP0911: <i>CP FIELD HOSPITALS (CPFH)</i>	5.333	3.435	1.929		1.929	3.498	1.539	0.000	0.000	Continuing	Continuing
• JP1111: <i>JOINT EXPEDITIONARY COLLECTIVE PROTECTION (JECP)</i>	0.000	0.000	0.000		0.000	0.000	4.714	20.863	44.885	Continuing	Continuing
	14.121	17.438	19.744		19.744	20.241	20.683	29.265	29.478	Continuing	Continuing

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C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011</u> <u>Base</u>	<u>FY 2011</u> <u>OCO</u>	<u>FY 2011</u> <u>Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To</u> <u>Complete</u>	<u>Total Cost</u>
• R12301: <i>CB PROTECTIVE SHELTER (CBPS)</i>											

D. Acquisition Strategy

JECP

Strategy based on evolutionary development in consonance with the JRO/User developed capability documents. During the Pre-MS A Concept Refinement Phase, conduct a tailored Analysis of Alternatives (AoA) leveraging the market survey, test results and lessons learned from the FY05 ColPro Technology Readiness Evaluation (TRE). During the Technology Development Phase following MS A, technology demonstrations were conducted to mitigate risk and identify affordable mature technologies that individually or together meet the Warfighters needs. Following MS B, a Statement of Work (SOW) and System Performance Specification (SPS) were used to award competitive cost plus incentive fee contract to build prototypes that will be subjected to robust engineering developmental testing and Operational Assessment during the System Development & Demonstration phase. Following MS C, award a Fixed Price Incentive Successive Target (FPIS) option for Low Rate Initial Production (LRIP) to support formal Developmental Testing (DT) and Multi-Service Operational Test & Evaluation (MOT&E). Following a successful Full Rate Production (FRP) decision, award a FPIS option with five one-year ordering periods. Full and open competition will be used with an updated SPS to award follow-on production contracts. Following JECP achieving Full Operational Capability, the Expeditionary Collective Protection-Enhanced Program will provide solutions to meet emerging and evolving User needs.

E. Performance Metrics

N/A

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Product Development (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** JECP - HW S - Prototype Development	C/CPIF	Science Applications International Corporation San Diego, CA	8.477	3.212	Jan 2010	1.330	Jan 2011	0.000		1.330	0.000	13.019	0.000
Subtotal			8.477	3.212		1.330		0.000		1.330	0.000	13.019	0.000

Remarks

Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** JECP - ES S - Systems Engineering IPT	MIPR	Various	3.234	1.061	Oct 2009	0.750	Oct 2010	0.000		0.750	0.000	5.045	0.000
ILS S - Integrated Logistics IPT	MIPR	Various	0.851	0.650	Oct 2009	0.500	Oct 2010	0.000		0.500	0.000	2.001	0.000
Subtotal			4.085	1.711		1.250		0.000		1.250	0.000	7.046	0.000

Remarks

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program										DATE: February 2010			
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>				R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>				PROJECT CO5: <i>COLLECTIVE PROTECTION (SDD)</i>					

Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** JECF - OTHT SB - Test & Evaluation IPT	MIPR	Various	3.645	0.940	Oct 2009	1.000	Oct 2010	0.000		1.000	0.000	5.585	0.000
OTHT SB - Prototype Performance Specification Testing	MIPR	Various	2.616	2.216	Oct 2009	0.000		0.000		0.000	0.000	4.832	0.000
DTE S - Prototype Production Qualification Testing	MIPR	Various	0.000	1.345	Jul 2010	9.310	Oct 2010	0.000		9.310	0.000	10.655	0.000
OTHT C - M98 Filter Set	MIPR	Various	0.000	0.000		2.350	Jan 2011	0.000		2.350	0.000	2.350	0.000
Subtotal			6.261	4.501		12.660		0.000		12.660	0.000	23.422	0.000

Remarks

Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** JECF - PM/MS S - APMO Support	MIPR	NSWC Dahlgren Dahlgren, VA	2.316	1.034	Oct 2009	1.000	Oct 2010	0.000		1.000	0.000	4.350	0.000
PM/MS S - APMO Contractor Support	C/FP	Solutions Development Corporation Dahlgren, VA	0.428	0.146	Jan 2010	0.132	Jan 2011	0.000		0.132	0.000	0.706	0.000

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Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
PM/MS S - JPM-ColPro Support	MIPR	NSWC Dahlgren Dahlgren, VA	0.673	0.570	Oct 2009	1.501	Oct 2010	0.000		1.501	0.000	2.744	0.000
PM/MS S - JPEO-CBD Support	MIPR	JPEO CBD Falls Church, VA	3.116	0.673	Oct 2009	0.586	Oct 2010	0.000		0.586	0.000	4.375	0.000
** ZSBIR - SBIR/STTR - Aggregated from ZSBIR-SBIR/STTR	MIPR	HQ AMC, Alexandria	0.000	0.155		0.000		0.000		0.000	0.000	0.155	0.000
Subtotal			6.533	2.578		3.219		0.000		3.219	0.000	12.330	0.000

Remarks

	Total Prior Years Cost	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
Project Cost Totals	25.356	12.002	18.459	0.000	18.459	0.000	55.817	0.000

Remarks

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT CO5: <i>COLLECTIVE PROTECTION (SDD)</i>

	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015							
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4				
** JECF - Prototype System Development & Testing	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■														
Operational Assessment (OA)													■	■	■	■	■															
Production Qualification Testing (PQT)									■	■	■	■	■	■	■	■	■															
Capability Production Document (CPD)																			■													
MS C Decision																			■													
LRIP Option																			■													
MOT&E																			■	■	■	■										
FRP Decision Review																															■	

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT CO5: <i>COLLECTIVE PROTECTION (SDD)</i>

Schedule Details

Event	Start		End	
	Quarter	Year	Quarter	Year
** JECF - Prototype System Development & Testing	4	2008	2	2013
Operational Assessment (OA)	2	2012	2	2013
Production Qualification Testing (PQT)	4	2010	1	2013
Capability Production Document (CPD)	2	2013	2	2013
MS C Decision	2	2013	2	2013
LRIP Option	2	2013	2	2013
MOT&E	2	2014	1	2015
FRP Decision Review	1	2015	1	2015

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT DE5: <i>DECONTAMINATION SYSTEMS (SDD)</i>
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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
DE5: <i>DECONTAMINATION SYSTEMS (SDD)</i>	16.611	36.786	28.499	0.000	28.499	23.944	25.770	14.701	5.928	Continuing	Continuing
Quantity of RDT&E Articles	0	14	13		13	0	0	0	0		

A. Mission Description and Budget Item Justification

This project funds System Development and Demonstration (SDD) for: (1) Decontamination Competitive Prototype; (2) the Decontamination Family of Systems (DFoS); (3) the Human Remains Decontamination System (HRDS); (4) Joint Platform Interior Decon (JPID); and (5) the Joint Service Sensitive Equipment Decontamination (JSSED).

The Decontamination Competitive Prototype (DC PROTO) effort will support the JSSED and JPID programs of record by performing risk mitigation and will identify a solution for the Joint Strike Fighter (JSF) peculiar interior/exterior decontamination requirement and to support their Live Fire testing in FY13. DC PROTO will evaluate prototype systems that will demonstrate the best decontamination technology for JSSED/JPID to increase sensitive equipment and platform interior decontamination capabilities. DC PROTO will evaluate other technologies that can be inserted into the JSSED/JPID programs to increase the capability of the selected JSSED/JPID technology while supporting the JSF test requirements.

The Decontamination Family of Systems (DFoS) program facilitates the rapid transition of mature Science & Technology research developments to existing JPM - Decon Programs of Record (PoR) and guides S&T community efforts toward meeting the needs of the Warfighter. Leveraging the outcomes of the Material Development Decision (MDD) (3rd Qtr FY10) directed Analysis of Alternatives, DFoS will develop a Family of Systems, to include equipment, to improve decontamination processes and decontaminant solutions to meet the capability gaps for decontaminating NTA and chemical and biological warfare agents from personnel, equipment, vehicle, ship, and aircraft interiors/exterior, terrain and fixed facility interiors/exterior.

The Human Remains Decontamination System (HRDS), Increment I, will utilize mature technologies to provide the capability for safe intra-theater handling and storage of Contaminated Human Remains (CHR) associated with a Chemical Warfare Agent (CWA) event. HRDS will be a Family-of-Systems (FoS) designed to leverage differing technology and requirements readiness across three systems: (1) a Contaminated Human Remains Pouch (CHRP) to support the initial recovery of CHR from Point of Fatality to a Mortuary Affairs Decontamination Collection Point (MADCP), (2) a Contaminated Remains Transfer Case System (CHRTS) capability to store or transport CHR post MADCP operations, and (3) a Remains Decontamination System (RDS) to support the capability to store or transport CHR post MADCP operations. The HRDS will provide the Services the capability to: 1) Safely recover, handle, and transport contaminated human remains prior to decontamination at a Mortuary Affairs Decontamination Control Point (MADCP); 2) Enable mortuary affairs units to safely perform their mission with a critical task being that of extensively

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT DE5: <i>DECONTAMINATION SYSTEMS (SDD)</i>
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documenting decedent data and obtaining DNA samples to facilitate positive identification of remains; 3) Fully decontaminate human remains (external), and; 4) Safely allow transport of decontaminated human remains from the MADCP to a final destination in the continental United States for final disposition.

The Joint Service Sensitive Equipment Decontamination System (JSSED) and Joint Platform Interior Decontamination (JPID) programs are based on a single technology and are being executed together by the Joint Material Decontamination System (JMDS) program office. These systems will fill the capability to decontaminate chemical and biological warfare agents from individual sensitive equipment, vehicle/aircraft/building interiors and the sensitive equipment within and the associated cargo. The JSSED will provide the first ever capability to decontaminate chemical and biological warfare agents from individual sensitive equipment that are high value or critical sensitive individual electronics and optics that cannot be decontaminated using existing methods without damage. The JPID will provide first ever capability to decontaminate chemical and biological warfare agents from platform interiors such as the interiors of vehicle/aircraft/building and the sensitive equipment within and the associated cargo. These capabilities allow the saving and reuse of contaminated critical and high value assets and avoid costly replacement of those assets. Neither of these capabilities currently exists in DoD.

Protective Self-Decontaminating Surfaces - CHRPS (Congressional Interest Item): Prototype field validation tests of VRCKappler Chemical Biohazard Protective systems. Lab test bacterial infections, deceases and contaminated human remains pouches (CHRP). Field and live test nerve gas and radiological agents.

Self Contained Automated Vehicle Washing Systems with Microwave Decontamination (Congressional Interest Item): Provide a large-vehicle washing and decontamination system with zero emissions to air or groundwater. Move from the prototype stage to full scale, portable decontamination wash down system.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) DC PROTO <i>FY 2010 Plans:</i> Conduct market survey/Industry Day/Sources Sought.	0.000	0.104	0.000	0.000	0.000
2) DC PROTO <i>FY 2010 Plans:</i> Select mature technologies capable of meeting JSSED and JPID requirements. Evaluate and test these technologies as compared against JSSED and JPID requirements. Conduct live agent efficacy and material compatibility testing.	0.000	8.657	0.000	0.000	0.000

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>		R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>		PROJECT DE5: <i>DECONTAMINATION SYSTEMS (SDD)</i>		
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
3) DC PROTO <i>FY 2011 Base Plans:</i> Continue material compatibility test and conduct early operational assessment and system integration evaluations on selected technologies for multiple airframes, tactical vehicles and sensitive equipment.		0.000	0.000	5.484	0.000	5.484
4) DFS <i>FY 2009 Accomplishments:</i> Conducted Industry Day/Request for Information/Evaluation of commercially available decontaminants (decon); developed prototype decon kit.		0.490	0.000	0.000	0.000	0.000
5) DFS <i>FY 2009 Accomplishments:</i> Standardized non-traditional agent (NTA) test methodology for decon; designed/developed/ demonstrated prototype environmentally controlled test fixture capability.		1.710	0.000	0.000	0.000	0.000
6) DFS <i>FY 2010 Plans:</i> Initiate development and evaluation test efforts for Surfactant System Technology, and Contamination Indicator/Decontamination Assurance Spray (DAS).		0.000	1.490	0.000	0.000	0.000
7) DFS <i>FY 2010 Plans:</i> Initiate the development effort to evaluate an aircraft decontaminant.		0.000	1.460	0.000	0.000	0.000
8) DFS		0.000	0.000	7.075	0.000	7.075

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT DE5: <i>DECONTAMINATION SYSTEMS (SDD)</i>				
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2011 Base Plans:</i> Conduct efficacy and deliver production representative articles/material compatibility testing/demonstrate manufacturing processes for Decon Wipes and eClO2.						
9) DFS <i>FY 2011 Base Plans:</i> Conduct testing and evaluation of Improved Reactive Sorbent/RSDL Reformulation/Identification-Assurance Sprays/Agent Disclosure Sprays and Protective Coatings.		0.000	0.000	1.436	0.000	1.436
10) DFS <i>FY 2011 Base Plans:</i> Conduct technology and manufacturing readiness assessments/initiate efficacy and material compatibility testing/transition Surfactant System and Contamination Indicator/DAS technologies to DFoS.		0.000	0.000	2.759	0.000	2.759
11) HRDS <i>FY 2010 Plans:</i> Conduct developmental testing and analysis of the Contaminated Human Remains Transfer Case (CHRT)		0.000	3.694	0.000	0.000	0.000
12) JPID <i>FY 2010 Plans:</i> Design and development of prototypes and logistics planning.		0.000	4.778	0.000	0.000	0.000
13) JPID		0.000	1.500	0.000	0.000	0.000

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program			DATE: February 2010			
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT DE5: <i>DECONTAMINATION SYSTEMS (SDD)</i>				
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
20) JSSED <i>FY 2011 Base Plans:</i> Complete Early Operational Assessment and conduct MS C.		0.000	0.000	2.054	0.000	2.054
21) JSSED <i>FY 2011 Base Plans:</i> Fabricate 4 JSSED Prototypes (at \$300 thousand each) for Operational Assessment testing.		0.000	0.000	1.200	0.000	1.200
22) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.		0.000	0.365	0.000	0.000	0.000
Accomplishments/Planned Programs Subtotals		16.611	28.422	28.499	0.000	28.499
		FY 2009	FY 2010			
Congressional Add: 1) Chemical and Biological Threat Reduction Coating <i>FY 2010 Plans:</i> Congressional Interest Item - Chemical and Biological Threat Reduction Coating		0.000	2.390			
Congressional Add: 2) Self-Decontaminating Polymer System for Chemical and Biological Warfare Agents. <i>FY 2010 Plans:</i> Congressional Interest Item - Self-Decontaminating Polymer System for Chemical and Biological Warfare Agents.		0.000	2.788			

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT DE5: <i>DECONTAMINATION SYSTEMS (SDD)</i>
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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010
Congressional Add: 3) Self Contained Automated Vehicle Washing Systems with microwave decontamination <i>FY 2010 Plans:</i> Congressional Interest Item - Self Contained Automated Vehicle Washing Systems with microwave decontamination.	0.000	1.593
Congressional Add: 4) Protective Self-Decontaminating Surfaces <i>FY 2010 Plans:</i> Congressional Interest Item - Protective Self-Decontaminating Surfaces - CHRPS.	0.000	1.593
Congressional Adds Subtotals	0.000	8.364

C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011</u> <u>Base</u>	<u>FY 2011</u> <u>OCO</u>	<u>FY 2011</u> <u>Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To</u> <u>Complete</u>	<u>Total Cost</u>
• JD0050: <i>DECONTAMINANT SYSTEM OF SYSTEMS</i>	0.000	0.000	0.000		0.000	3.280	4.468	6.884	7.029	Continuing	Continuing
• JD0055: <i>JOINT SERVICE PERSONNEL/SKIN DECON SYSTEM (JSPDS)</i>	8.280	4.466	0.000		0.000	0.000	0.000	8.645	9.105	Continuing	Continuing
• JD0056: <i>JS TRANS DECON SYSTEM - SMALL SCALE (JSTDS-SS)</i>	12.124	21.940	18.160		18.160	12.924	7.900	5.455	4.459	Continuing	Continuing
• JD0060: <i>JOINT PLATFORM INTERIOR DECON (JPID)</i>	0.000	0.000	0.000		0.000	4.097	14.064	18.977	25.604	Continuing	Continuing
	0.000	0.000	0.000		0.000	14.648	0.000	0.000	0.000	Continuing	Continuing

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT DE5: <i>DECONTAMINATION SYSTEMS (SDD)</i>
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C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011</u> <u>Base</u>	<u>FY 2011</u> <u>OCO</u>	<u>FY 2011</u> <u>Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To</u> <u>Complete</u>	<u>Total Cost</u>
• JD0061: <i>JS SENSITIVE EQUIP DECON (JSSED)</i>											

D. Acquisition Strategy

DC PROTO

DC PROTO will conduct a Sources Sought for a prototype suitable for sensitive equipment and platform interior decontamination prototypes. The competitive prototype results will be integrated into the JSSED and JPID programs for program risk reduction.

DFS

DFoS will utilize an incremental acquisition strategy to transition various developmental technology efforts (COTS, JSTO, DTRA efforts, etc.) to fill current and future capability gaps. DFoS will support MDAPs and Programs of Record (POR) by guiding S&T efforts and transitioning mature technologies to meet program requirements. The DFoS acquisition will be managed as a Family-of-Systems (FoS), leveraging differing technologies in each subsystem to fulfill Warfighter capability gaps. A multi-phased Analysis of Alternatives (AoA) will be conducted to identify and evaluate the operational effectiveness of potential material solutions to satisfy Service requirements. As each AoA phase is completed, individual systems and their respective phases of entry will be identified. Industry will be solicited and through competitive prototyping, material solutions will be down-selected for continued development and fielding as a new joint force capability.

HRDS

The Human Remains Decontamination System (HRDS) acquisition will be managed as a Family-of-Systems (FoS), leveraging differing technologies in each subsystem to fulfill Warfighter capability gaps. A multi-phased Analysis of Alternatives (AoA) is being conducted for the HRDS FoS to identify and evaluate the operational effectiveness of potential material solutions to satisfy Service requirements. As each AoA phase is completed, individual systems and their respective phases of entry will be identified. Industry will be solicited and through competitive prototyping, material solutions will be down-selected for continued development and fielding as a new joint force capability.

JSSED

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT DE5: <i>DECONTAMINATION SYSTEMS (SDD)</i>
<p>The Joint Service Sensitive Equipment Decontamination (JSSED) programs will be acquired as part of the overarching Joint Material Decontamination System (JMDS) evolutionary acquisition strategy that covers both the Joint Platform Interior Decontamination (JPID) and the JSSED. This strategy will use a single technology to meet the individual sensitive equipment through incremental development. The JSSED strategies is under the JMDS contracting strategy that awarded one single base Engineering and Manufacturing Development (EMD) contract (Cost Plus Incentive Fee) with Low Rate Initial Production and Full Rate Production options (Fixed Price Successive Target) in open competition. The JMDS program will integrate the competitive prototype effort into the JMDS Milestone C/LRIP Decision.</p>		
E. Performance Metrics N/A		

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT DE5: <i>DECONTAMINATION SYSTEMS (SDD)</i>
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Product Development (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** CONG - Congressional Interest Item - HW C - Chemical and Biological Threat Reduction Coating	C/CPFF	TBD	0.000	2.390	Apr 2010	0.000		0.000		0.000	0.000	2.390	0.000
Congressional Interest Item - HW C - Self Decontaminating Polymer System for Chemical and Biological Warfare Agents	C/CPFF	TBD	0.000	2.788	Apr 2010	0.000		0.000		0.000	0.000	2.788	0.000
** DFS - HW S - Decon wipes	MIPR	RDECOM-Natick MA	0.000	0.000		1.792	Jan 2011	0.000		1.792	0.000	1.792	0.000
HW S - Electro-chemically Generated Chlorine Dioxide (eClO2)	MIPR	RDECOM-Natick MA	0.000	0.000		2.000	Jan 2011	0.000		2.000	0.000	2.000	0.000
HW C - Surfactant System	MIPR	Defense Threat Reduction Agency (DTRA) Ft. Belvoir, VA	0.000	0.188	Oct 2009	0.000		0.000		0.000	0.000	0.188	0.000
HW C - Contaminant Indicator/ Decon Assurance Spray	MIPR	Defense Threat Reduction Agency (DTRA) Ft. Belvoir, VA	0.000	0.200	Oct 2009	0.600	Apr 2011	0.000		0.600	0.000	0.800	0.000
HW C - Aircraft Decon	MIPR	RDECOM-Natick MA	0.000	0.945	Jan 2010	0.000		0.000		0.000	0.000	0.945	0.000
	C/CPFF	TBD	0.000	1.593	Apr 2010	0.000		0.000		0.000	0.000	1.593	0.000

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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Product Development (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
Congressional Interest Item - HW C - Self Contained Automated Vehicle Washing Systems with microwave decontamination													
** HRDS - Congressional Interest Item - HW C - Protective Self-Decontaminating Surfaces - CHRPS	C/CPFF	TBD	0.000	1.593	Apr 2010	0.000		0.000		0.000	0.000	1.593	0.000
** JPID - HW C - SDD Contract, EMD Contract and fabrication	C/CPIF	Teledyne Brown Engineering Huntsville, AL	0.000	3.000	Jan 2010	2.815	Jan 2011	0.000		2.815	0.000	5.815	0.000
** JSSED - HW S - EMD Contract - System Development and Fabrication	C/CPIF	Teledyne Brown Engineering - Huntsville AL	15.764	3.000	Jan 2010	0.500	Jan 2011	0.000		0.500	0.000	19.264	0.000
Subtotal			15.764	15.697		7.707		0.000		7.707	0.000	39.168	0.000

Remarks

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT DE5: <i>DECONTAMINATION SYSTEMS (SDD)</i>
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Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** DC PROTO - Market Survey/Sources Sought Assessment	MIPR	RDECOM Natick, MA	0.000	0.104	Oct 2009	0.000		0.000		0.000	0.000	0.104	0.000
** DFS - ES S - IPT Technical Support	MIPR	Various	0.226	0.200	Jan 2010	1.124	Jan 2011	0.000		1.124	0.687	2.237	0.000
** HRDS - TD/D SB - SME Technical Support	MIPR	TBD	0.000	0.643	Jan 2010	0.000		0.000		0.000	0.000	0.643	0.000
Subtotal			0.226	0.947		1.124		0.000		1.124	0.687	2.984	0.000

Remarks

Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** DC PROTO - Competitive Prototype Testing	MIPR	TBD	0.000	8.144	Apr 2010	4.892	Oct 2010	0.000		4.892	0.000	13.036	0.000
** DFS - DTE S - Electro-Chemically Generated Chlorine Dioxide (eClO2)	MIPR	TBD	0.000	0.000		1.134	Jan 2011	0.000		1.134	0.221	1.355	0.000
DTE S - Decon Wipes	MIPR	TBD	0.000	0.000		0.981	Jan 2011	0.000		0.981	0.165	1.146	0.000

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT DE5: <i>DECONTAMINATION SYSTEMS (SDD)</i>
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Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
OTHT SB - Improved Reactive Sorbent	MIPR	TBD	0.000	0.000		0.571	Oct 2010	0.000		0.571	0.000	0.571	0.000
DTE C - Surfactant System	MIPR	Army Evaluation Center Alexandria, VA	0.000	0.188	Jan 2010	0.650	Jan 2011	0.000		0.650	0.094	0.932	0.000
DTE C - Contaminant Indicator/Decon Assurance Spray	MIPR	TBD	0.000	0.200	Apr 2010	0.875	Apr 2011	0.000		0.875	0.188	1.263	0.000
DTE C - Reactive Skin Decontamination Lotion Reformulation	MIPR	TBD	0.000	0.000		0.225	Apr 2011	0.000		0.225	0.000	0.225	0.000
** HRDS - OTHT SB - Prototype Planning and Testing	MIPR	TBD	0.000	1.390	Apr 2010	0.000		0.000		0.000	0.000	1.390	0.000
** JPID - JPID Development Testing	MIPR	ATEC Aberdeen Proving Ground, MD	0.000	2.478	Oct 2009	3.960	Oct 2010	0.000		3.960	0.000	6.438	0.000
** JSSED - OTHT SB - JSSED/JMDS developmental test planning/execution	MIPR	ATEC Aberdeen, MD	1.244	1.500	Oct 2009	2.062	Oct 2010	0.000		2.062	0.000	4.806	0.000
Subtotal			1.244	13.900		15.350		0.000		15.350	0.668	31.162	0.000

Remarks

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** DC PROTO - Program Management Support	MIPR	RDECOM Natick, MA	0.000	0.513	Oct 2009	0.592	Oct 2010	0.000		0.592	0.000	1.105	0.000
** DFS - PM/MS S - DFoS Integrated Product Team Support	MIPR	RDECOM-Natick MA	0.870	0.674	Oct 2009	0.865	Oct 2010	0.000		0.865	0.000	2.409	0.000
PM/MS S - Program Support	MIPR	Marine Corps Systems Command Quantico, VA	0.265	0.355	Jan 2010	0.453	Jan 2011	0.000		0.453	0.000	1.073	0.000
** HRDS - PM/MS SB - Program Office Support	MIPR	RDECOM-Natick MA	0.000	1.661	Jan 2010	0.000		0.000		0.000	0.000	1.661	0.000
** JPID - JPID Service Integrated Product Team Support	MIPR	Various	0.000	0.800	Oct 2009	1.716	Oct 2010	0.000		1.716	0.000	2.516	0.000
** JSSED - PM/MS S - JSSED/JMDS Service Integrated Product Team Support	MIPR	Various	6.673	1.874	Oct 2009	0.692	Oct 2010	0.000		0.692	0.000	9.239	0.000
** ZSBIR - SBIR/STTR - Aggregated from ZSBIR-SBIR/STTR	MIPR	HQ AMC, Alexandria	0.000	0.365		0.000		0.000		0.000	0.000	0.365	0.000
Subtotal			7.808	6.242		4.318		0.000		4.318	0.000	18.368	0.000

Remarks

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
** CONG - Self contained automated vehicle washing systems with microwave decontamination							■	■	■	■	■	■																
Protective Self-Decontaminating Surfaces - CHRPS							■	■	■	■	■	■																
** DC PROTO - Market Survey/Industry Day/ Sources Sought						■	■																					
Competitive Prototype Test							■	■	■	■	■	■																
** DFS - Commercial Decontaminant	■	■	■																									
NTA Decon Assessment Capability				■																								
Decon Wipes					■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■				
Electro-chemically Generated Chlorine Dioxide (eClO2)					■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■				
Surfactant System					■	■	■	■	■	■	■	■	■	■	■	■												
Contaminant Indicator/Decon Assurance Spray					■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■								
Aircraft Decon							■	■	■	■	■	■	■	■	■	■												
** HRDS - CHRT Market Survey	■																											
HRDS MDD				■																								
HRDS Document Preparation, technical support, and test planning					■	■	■	■	■	■																		
CHRP/CHRT Development Testing									■	■																		

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT DE5: <i>DECONTAMINATION SYSTEMS (SDD)</i>

	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015				
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	
CHRP/CHRT MS C												■	■																
CHRP/CHRT Fielding													■	■	■	■	■	■	■	■	■	■	■						
** JPID - JPID Systems Design and Development	■	■	■	■																									
JPID Developmental Test			■	■	■	■	■	■	■	■	■	■																	
JPID Early Operational Assessment								■	■																				
JPID Competitive Prototype							■	■	■	■	■																		
JPID Milestone C LRIP																■													
JPID MOT&E													■	■	■	■	■												
JPID MS/C FRP Decision																				■									
JPID Full Rate Production																				■	■	■	■	■	■	■	■	■	
** JSSED - JSSED/JMDS System Development	■	■	■	■																									
JSSED/JMDS Developmental Test			■	■	■	■	■	■																					
JSSED/JMDS Early Operational Assessment								■	■																				
JSSED/JMDS Competitive Prototype							■	■	■	■	■																		
JSSED/JMDS MS C & LRIP DECISION																■													
JSSED/JMDS Early Operational Assessment #2								■	■																				
JSSED/JMDS MOT&E													■	■	■	■	■												
																				■									

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT DE5: <i>DECONTAMINATION SYSTEMS (SDD)</i>
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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
JSSSED/JMDS Full Rate Production (FRP) DECISION																												
JSSSED/JMDS FRP																	■	■	■	■	■	■	■	■	■	■	■	■

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
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Schedule Details

Event	Start		End	
	Quarter	Year	Quarter	Year
** CONG - Self contained automated vehicle washing systems with microwave decontamination	3	2010	4	2011
Protective Self-Decontaminating Surfaces - CHRPS	3	2010	4	2011
** DC PROTO - Market Survey/Industry Day/Sources Sought	1	2010	2	2010
Competitive Prototype Test	3	2010	4	2011
** DFS - Commercial Decontaminant	1	2009	3	2009
NTA Decon Assessment Capability	4	2009	4	2009
Decon Wipes	2	2010	4	2014
Electro-chemically Generated Chlorine Dioxide (eClO2)	2	2010	4	2014
Surfactant System	2	2010	2	2013
Contaminant Indicator/Decon Assurance Spray	2	2010	1	2014
Aircraft Decon	3	2010	2	2013
** HRDS - CHRT Market Survey	1	2009	1	2009
HRDS MDD	4	2009	4	2009
HRDS Document Preparation, technical support, and test planning	2	2010	2	2011
CHRP/CHRT Development Testing	1	2011	2	2011
CHRP/CHRT MS C	3	2011	4	2011
CHRP/CHRT Fielding	4	2011	2	2014
** JPID - JPID Systems Design and Development	1	2008	4	2009

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
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Event	Start		End	
	Quarter	Year	Quarter	Year
JPID Developmental Test	3	2009	4	2011
JPID Early Operational Assessment	4	2010	1	2011
JPID Competitive Prototype	3	2010	3	2011
JPID Milestone C LRIP	4	2011	4	2011
JPID MOT&E	1	2012	1	2013
JPID MS/C FRP Decision	3	2013	3	2013
JPID Full Rate Production	3	2013	4	2015
** JSSED - JSSED/JMDS System Development	1	2008	4	2009
JSSED/JMDS Developmental Test	3	2009	4	2010
JSSED/JMDS Early Operational Assessment	4	2010	1	2011
JSSED/JMDS Competitive Prototype	3	2010	3	2011
JSSED/JMDS MS C & LRIP DECISION	4	2011	4	2011
JSSED/JMDS Early Operational Assessment #2	4	2010	1	2011
JSSED/JMDS MOT&E	1	2012	1	2013
JSSED/JMDS Full Rate Production (FRP) DECISION	3	2013	3	2013
JSSED/JMDS FRP	3	2013	4	2015

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program								DATE: February 2010			
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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
IP5: <i>INDIVIDUAL PROTECTION (SDD)</i>	18.363	21.094	9.678	0.000	9.678	4.833	3.044	0.756	0.563	Continuing	Continuing
Quantity of RDT&E Articles	0	544	0		0	0	0	0	0		

A. Mission Description and Budget Item Justification

This project funds System Development and Demonstration (SDD) of individual protection equipment, the goal is to provide equipment that allows the individual soldier, sailor, airman, or marine to operate in a contaminated Nuclear, Biological and Chemical (NBC) environment with little or no degradation of his/her performance.

The three efforts listed below are funded in this program:

(1) The Joint Service Aircrew Mask (JSAM) is an Acquisition Category (ACAT) III Family of Systems (FoS) respiration system being incrementally developed. JSAM MPU-6 Apache is for use with the Integrated Helmet And Display Sighting System, JSAM Fixed Wing (FW) MBU-25/26 respirator and JSAM MPU-5 Rotor Wing (RW) are being developed for use in the majority of the Department of Defense's (DoD's) Fixed and Rotary Wing aircraft. The F-35 JSAM MBU-26 is being developed with the FW JSAM MBU-25 to meet the needs of the Major Defense Acquisition Program, the Joint Strike Fighter (JSF). The goal of the overall JSAM project is to develop, manufacture, field and sustain an aircrew respirator system that, in conjunction with a below-the-neck (BTN) clothing ensemble, will provide the capability for all aircrew to fly throughout their full operating envelope in an actual or perceived Chemical and Biological (CB) warfare environment. JSAM will be a lightweight CB protective mask that will be worn as CB protection for most Army, Air Force, Navy and Marine rotary and fixed-wing aircrew members. The FW JSAM will be the first and only CB protective mask in the DoD inventory that can provide anti-G protection, up to 9 times the vertical force (Gz), for aircrew in high performance aircraft. All JSAM Increments will be compatible with most below-the-neck CB ensembles and existing aircrew life support equipment. They will include a protective hood assembly, CB filter, blower assembly, and an intercom for ground communication. They will provide flame and thermal protection, provide hypoxia protection to 60,000 feet, demist/emergency demist and anti-drown features. The MPU-5 and MPU-6 variants are being designed to be capable of being donned/doffed in flight.

(2) The Joint Service General Purpose Mask (JSGPM) funds SDD of respiratory and ocular protection technologies aimed at providing incremental upgrades for the JSGPM. Additionally, this project funds the Technology Development (TD) phase of the Advanced Respiratory Protection Initiative (ARPI) program for developing revolutionary materials, design and concepts that may be transitioned into future Chem/Bio ensemble (Joint Chemical Ensemble). Performance enhancements for all respiratory and ocular protection programs will be focused on increasing the protection levels of the systems from Chemical Warfare Agents (CWAs) and Toxic Industrial Chemicals (TICs) while reducing the physiological and logistical burdens.

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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(3) The Lightweight Chemical Biological Ensemble (LCBE), aims to provide CBRN individual protection for the Warfighter while reducing physiological and logistical burdens.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p>1) JSAM</p> <p><i>FY 2009 Accomplishments:</i> JSAM MPU-6 (Apache) - Completed Operational Testing (OT). Prepared documentation for Milestone (MS) C Full Rate Production (FRP). JSAM MPU-5 (RW) - Continued Developmental Testing (DT). JSAM FW - Initiated flight testing, Chem/Bio, environmental and continued integration testing for joint service aircraft platform.</p> <p><i>FY 2010 Plans:</i> JSAM MPU-5 (RW) - Prepare specific mask tooling for prototypes. Complete DT. Produce MPU-5 prototypes (256 units at a cost of \$4,400 ea) for OT. Start OT. JSAM FW - Continue and complete DT flight testing. Start OT. Produce prototypes (288 prototype at a unit cost of \$4,130 ea) for OT. Start OT.</p> <p><i>FY 2011 Base Plans:</i> JSAM MPU-5 (RW) - Complete OT. Prepare documentation for MS C FRP. JSAM FW - Complete OT. Prepare documentation for MS C FRP.</p>	16.781	14.714	7.269	0.000	7.269
<p>2) JSGPM</p> <p><i>FY 2010 Plans:</i> JSGPM (ARPI) - Conduct government testing screening. Initiate filter qualification testing on potential End of Service Life Indicator (ESLI) candidates.</p>	0.000	1.444	2.409	0.000	2.409

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010				
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B. Accomplishments/Planned Program (\$ in Millions)								
				FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2011 Base Plans:</i> JSGPM (ARPI) - Conduct government testing to ensure carbons transitioned to JSGPM filters to improve TIC protection meeting the user requirements. Conduct government testing on novel filtration candidates considered for Joint Chemical Ensemble (JCE).</p> <p>JSGPM - Complete testing of ESLI.</p>								
3) LCBE				0.000	2.305	0.000	0.000	0.000
<p><i>FY 2010 Plans:</i> Prepare MS A documentation for LCBE Increment 1. Technology demonstration of lightweight garment technologies and designs. Continue thermal burden reduction/heat stress assessment.</p>								
4) SBIR				0.000	0.241	0.000	0.000	0.000
<p><i>FY 2010 Plans:</i> Small Business Innovative Research.</p>								
Accomplishments/Planned Programs Subtotals				16.781	18.704	9.678	0.000	9.678
				FY 2009	FY 2010			
Congressional Add: 1) JSAM				1.582	2.390			
<p><i>FY 2009 Accomplishments:</i> Congressional Interest Item - JSAM Donn\Doff. Developed Donn\Doff capability for MPU-5 and MPU-6 requirement.</p>								

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010
<i>FY 2010 Plans:</i> Congressional Interest Item - JSAM Donn\Doff. Continue development of Donn\Doff capability for MPU-5 and MPU-6 requirement.		
Congressional Adds Subtotals	1.582	2.390

C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011</u> <u>Base</u>	<u>FY 2011</u> <u>OCO</u>	<u>FY 2011</u> <u>Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To</u> <u>Complete</u>	<u>Total Cost</u>
• IP7: <i>INDIVIDUAL PROTECTION OPERATIONAL SYS DEV</i>	4.560	0.000	0.000		0.000	2.869	4.371	7.548	4.681	Continuing	Continuing
• JI0002: <i>JS AIRCREW MASK (JSAM)</i>	0.000	23.045	6.964		6.964	12.919	12.112	14.084	9.017	Continuing	Continuing
• JI0003: <i>JOINT SERVICE GENERAL PURPOSE MASK (JSGPM/JSCESM)</i>	42.391	48.282	49.835		49.835	51.508	56.463	56.334	60.975	Continuing	Continuing
• JI0300: <i>JOINT CHEMICAL ENSEMBLE (JCE)</i>	0.000	0.000	0.000		0.000	0.000	7.446	9.918	6.936	Continuing	Continuing
• MA0400: <i>PROTECTIVE CLOTHING (JSLIST)</i>	37.484	20.393	17.887		17.887	18.208	9.429	6.943	6.944	Continuing	Continuing

D. Acquisition Strategy

JSAM

The JSAM Acquisition Program Baseline Agreement (APBA) identifies JSAM Type IA Apache (MPU-6) as the Rotary Wing (RW) Integrated Helmet and Display Sighting System (IHADSS) variant. The JSAM Type I RW (MPU-5) that is being developed for the majority of RW aircrew. JSAM Type IA Apache (MPU-6) will be fielded first. Appropriate production options will be exercised.

The JSAM Type II Fixed Wing (FW) variant will meet the needs of the FW aircrew, and majority of the requirements for the JSF JSAM.

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
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<p>JSGPM</p> <p>JSGPM: All possible candidates will be identified through the Request For Information (RFI). The candidates will be screened against CWAs and TICs at the sorbent level. Candidates that show an indication that it may provide a performance enhancement may be transitioned into filter qualification testing. The qualification of a new filtration media for JSGPM will be based on the current JSGPM filter specification.</p> <p>JSGPM (ARPI): The Advanced Respiratory Protection Initiative (ARPI) program will be based on full and open competition. A Request For Information was released in July 2008 to evaluate what novel concepts, materials and designs that could be pursued for the next generation system. An analysis of the results of the market survey will be conducted and potential candidates will be pursued for further evaluation.</p> <p>LCBE</p> <p>The Lightweight Chemical Biological Ensemble (LCBE) program will pursue an evolutionary incremental approach to provide capability to the Warfighter. Each increment of LCBE will provide technologies with military utility that are modular in function, and offer improvement in form and fit over current systems. The LCBE program will develop, integrate, test, procure and field systems that increase Warfighter operational performance in a CBRN environment via the use of emerging technologies and by leveraging tradespace in areas such as protection level, heat stress, durability, antimicrobial properties, launderability, self-detoxification, protection time, etc. Where appropriate, modeling and simulation tools will be used to lower LCBE program risks, reduce costs and ensure a high confidence in selected technologies.</p> <p>LCBE INCREMENT 1</p> <p>The LCBE will use an evolutionary acquisition strategy with phased development. The first LCBE increment will provide an operationally useful and supportable capability in as short a time as possible. Accordingly, Increment 1 of LCBE will incorporate an accelerated development cycle leveraging existing COTS technologies that will, at a minimum, provide a lightweight CB protective garment capability. Gate testing and down-selection of prototypes will comprise the initial phases of the Government's testing program. A competitively awarded contract is planned for DT and Operational Assessment (OA) will occur prior to MS C. Appropriate system requirements reviews, test readiness reviews, producibility reviews and audits will be scheduled as required prior to each milestone.</p> <p>Future increments of LCBE shall be defined via separate Capability Development Document (CDDs)/Capability Production Document (CPDs) and will follow a similar path/process from MS A or MS B through MS C/FRP and will leverage preceding efforts to the greatest extent possible, maintaining commonality and synergy across all increments.</p>		

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY	R-1 ITEM NOMENCLATURE	PROJECT
0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	IP5: <i>INDIVIDUAL PROTECTION (SDD)</i>

E. Performance Metrics

N/A

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT IP5: <i>INDIVIDUAL PROTECTION (SDD)</i>
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Product Development (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** JSAM - HW S - Contractor Development Types I/IA	C/CPAF	AVOX Lancaster, NY	36.353	0.795	Oct 2009	0.000		0.000		0.000	0.000	37.148	7.209
SW SB - Contractor Development Type II	C/FPI	Gentex Rancho Cucamonga, CA	12.658	4.626	Oct 2009	0.425	Jan 2011	0.000		0.425	0.000	17.709	0.000
HW S - Donn/Doff development	C/FFP	Gentex Rancho Cucamonga, CA	1.582	1.625	Apr 2010	0.000		0.000		0.000	0.000	3.207	0.000
Subtotal			50.593	7.046		0.425		0.000		0.425	0.000	58.064	7.209

Remarks

Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** JSGPM - ES C - JSGPM Filter	MIPR	ECBC APG, MD	0.000	0.118	Oct 2009	0.215	Oct 2010	0.000		0.215	0.000	0.333	0.000
ES C - JSGPM Filter	MIPR	NRL Washington, DC	0.000	0.100	Oct 2009	0.150	Oct 2010	0.000		0.150	0.000	0.250	0.000
** LCBE - ES S - Engineering IPT	MIPR	Various	0.000	0.400	Apr 2010	0.000		0.000		0.000	0.000	0.400	0.000

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Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
ES SB - Tech Demos	MIPR	Various	0.000	1.360	Oct 2009	0.000		0.000		0.000	0.000	1.360	0.000
Subtotal			0.000	1.978		0.365		0.000		0.365	0.000	2.343	0.000

Remarks

Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** JSAM - OTHS SB - Govt Dev Test	MIPR	Various	24.347	2.294	Oct 2009	0.000		0.000		0.000	7.174	33.815	0.092
OTE S - Govt Operational Test Type II	MIPR	Various	9.706	5.064	Oct 2009	4.049	Jan 2011	0.000		4.049	4.046	22.865	0.404
OTHT SB - Govt Operational Test Type I	C/FFP	AVOX Lancaster, NY	3.074	0.000		1.980	Oct 2010	0.000		1.980	0.000	5.054	0.185
** JSGPM - DTE SB - JSGPM Filter Testing	MIPR	Various	0.000	0.776	Oct 2009	1.594	Oct 2010	0.000		1.594	0.000	2.370	0.000
DTE SB - JSGPM Filter Testing	MIPR	NRL Washington, DC	0.000	0.250	Oct 2009	0.250	Oct 2010	0.000		0.250	0.000	0.500	0.000
Subtotal			37.127	8.384		7.873		0.000		7.873	11.220	64.604	0.681

Remarks

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** JSAM - PM/MS C - Program Management/ Management Support	MIPR	Various	17.525	2.700	Oct 2009	0.815	Oct 2010	0.000		0.815	0.000	21.040	5.421
** JSGPM - PM/MS C - Conduct Market Survey Analysis	MIPR	JPMO IP Stafford, VA	0.000	0.200	Oct 2009	0.200	Oct 2010	0.000		0.200	0.000	0.400	0.000
** LCBE - PM/MS S - JPM Support	MIPR	Various	0.000	0.545	Apr 2010	0.000		0.000		0.000	0.000	0.545	0.000
** ZSBIR - SBIR/STTR - Aggregated from ZSBIR-SBIR/STTR	MIPR	HQ AMC, Alexandria	0.000	0.241		0.000		0.000		0.000	0.000	0.241	0.000
Subtotal			17.525	3.686		1.015		0.000		1.015	0.000	22.226	5.421

Remarks

Project Cost Totals	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
Project Cost Totals	105.245	21.094		9.678		0.000		9.678	11.220	147.237	13.311

Remarks

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
** JSAM - DT MPU-5 Apache	■																											
OT&E MPU-5 Apache	■	■	■																									
MS C FRP Decision MPU-5 Apache			■	■																								
IOC MPU-5 Apache						■	■																					
FOC MPU-5 Apache													■															
DT MPU-6 RW	■	■	■	■	■																							
MS C LRIP Decision MPU-6 Rotor Wing							■																					
OT&E MPU-6 RW											■	■																
MS C FRP MPU-6 Rotor Wing											■																	
IOC MPU-6 RW																	■											
DT MBU-25/26 FW	■	■	■	■	■																							
Milestone C (LRIP) MBU-25/26 FW									■																			
OT&E MBU-25/26 FW												■	■															
MS C FRP Decision MBU-25/26 FW																■												
IOC MBU-25 /26 FW																	■											
** JSGPM - JSGPM Sorbent Testing						■	■																					
JSGPM Filter Qualification Testing							■	■	■																			
JSGPM Fielding Decision											■																	
JSGPM (ARPI) Market Survey Analysis						■	■																					
JSGPM (ARPI) Method Verification							■	■																				

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT IP5: <i>INDIVIDUAL PROTECTION (SDD)</i>

	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015				
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	
JSGPM (ARPI) Candidate Screening							■	■	■	■	■																		
JSGPM (ARPI) Down-Select												■																	
JSGPM (ARPI) Advanced Design Transition Assessments									■	■	■	■																	
JSGPM (ARPI) Integration Testing													■	■	■	■													
** LCBE - ESLI Test & Evaluation		■	■	■	■																								
LCBE Start IPT								■																					
LCBE Start OT														■	■														
LCBE MS C														■															

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT IP5: <i>INDIVIDUAL PROTECTION (SDD)</i>
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Schedule Details

Event	Start		End	
	Quarter	Year	Quarter	Year
** JSAM - DT MPU-5 Apache	4	2007	1	2009
OT&E MPU-5 Apache	2	2008	3	2009
MS C FRP Decision MPU-5 Apache	3	2009	4	2009
IOC MPU-5 Apache	2	2010	3	2010
FOC MPU-5 Apache	2	2012	2	2012
DT MPU-6 RW	4	2007	1	2010
MS C LRIP Decision MPU-6 Rotor Wing	3	2010	3	2010
OT&E MPU-6 RW	2	2011	3	2011
MS C FRP MPU-6 Rotor Wing	3	2011	3	2011
IOC MPU-6 RW	1	2013	1	2013
DT MBU-25/26 FW	1	2008	1	2010
Milestone C (LRIP) MBU-25/26 FW	4	2010	4	2010
OT&E MBU-25/26 FW	3	2011	4	2011
MS C FRP Decision MBU-25/26 FW	2	2012	2	2012
IOC MBU-25 /26 FW	4	2012	4	2012
** JSGPM - JSGPM Sorbent Testing	1	2010	2	2010
JSGPM Filter Qualification Testing	3	2010	1	2011
JSGPM Fielding Decision	2	2011	2	2011

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Event	Start		End	
	Quarter	Year	Quarter	Year
JSGPM (ARPI) Market Survey Analysis	1	2010	2	2010
JSGPM (ARPI) Method Verification	3	2010	4	2010
JSGPM (ARPI) Candidate Screening	3	2010	3	2011
JSGPM (ARPI) Down-Select	4	2011	4	2011
JSGPM (ARPI) Advanced Design Transition Assessments	1	2011	4	2011
JSGPM (ARPI) Integration Testing	1	2012	4	2012
** LCBE - ESLI Test & Evaluation	2	2009	1	2010
LCBE Start IPT	4	2010	4	2010
LCBE Start OT	2	2012	3	2012
LCBE MS C	2	2012	2	2012

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program								DATE: February 2010			
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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
IS5: <i>INFORMATION SYSTEMS (SDD)</i>	45.694	27.301	13.844	0.000	13.844	24.984	24.872	25.345	25.775	Continuing	Continuing
Quantity of RDT&E Articles	80	0	0		0	0	0	0	0		

A. Mission Description and Budget Item Justification

This funding supports System Development and Demonstration and Low Rate Initial Production (SDD/LRIP).

Efforts funded in this project are: (1) Joint Effects Model (JEM); (2) Joint Operational Effects Federation (JOEF); (3) the Joint Warning and Reporting Network (JWARN); and (4) the JPEO-CBD Software Support Activity (SSA).

The JEM is DoD's only accredited model for predicting hazards associated with the release of contaminants into the environment. JEM is being developed in separate increments and is capable of modeling hazards in a variety of scenarios including: counterforce, passive defense, accident and/or incidents (Increment 1); high altitude releases, urban NBC environments (Increment 2); building interiors, and human performance degradation (Increment 3). Battle space commanders and first responders must have a Chemical, Biological, Radiological, Nuclear (CBRN) hazard prediction capability in order to make decisions that will minimize risks of CBRN contamination and enable them to continue mission operations. JEM operates in an integrated fashion with operational and tactical Command, Control, Communications, Computers, Intelligence, Surveillance and Reconnaissance (C4ISR) systems, and in a standalone mode. JEM will interface and communicate with the other programs such as JWARN, JOEF, weather systems, intelligence systems, and various databases.

JOEF will be a near real-time course of action analysis tool developed in three increments using a detailed NBC hazard prediction model. Each increment supports Aerial Ports of Debarkation (APODs), Sea Ports of Debarkation (SPODs), mobile forces, medical and automated Tactics, Techniques and Procedures (TTPs) in various levels of fidelity. Increment 1 will support deliberate planning for operational and strategic users in a C4ISR common operating environment (COE); Command and Control Personal Computers (C2PC); and crisis planning for the operational users in a COE.

The Joint Warning and Reporting Network (JWARN) will provide, in the first of two increments, joint forces with a comprehensive analysis and response capability to minimize the effects of hostile CBRN attacks, as well as, accidents and incidents. It will provide the operational capability to employ NBC warning technology which will collect, analyze, identify, locate, report, and disseminate NBC warnings. JWARN will be compatible and integrated with Joint Services C4ISR Systems. JWARN is transition from COE standards to Service Oriented Architecture (SOA). JWARN Increment 2 will provide an expansion of sensors that will connect to JWARN, increased automation of message handling, improved false alarm filtering, integration of route-planning calculator, and interoperability with additional C2 systems.

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JWARN will be located in Command and Control Centers at the appropriate level and will be employed by CBRN defense specialists and other designated personnel. This employment will transfer data automatically from existing sensors and to and from the future sensors to provide commanders with the capability to support operational decision making in a CBRN environment. JWARN will provide additional data processing to support the production of plans and reports, and access to specific CBRN information to improve the efficiency of limited CBRN personnel assets. JWARN will integrate existing sensors into a sensor network or host C2 system, but does not provide the sensors that will be employed in the operating environment.

The JPEO-CBD SSA is a JPEO-CBD enterprise-wide, user developmental support and service organization focusing on development assistance and net-centric interoperability. The SSA provides the CBRN Warfighter with Joint Service solutions for Integrated Architectures, Information Assurance, Verification, Validation and Accreditation (VV&A) and Data Management; interoperable and integrated net-centric, Service-oriented, composable solutions for CBD; and infusion of latest technologies into programs of record. CBRN user community and related communities of interest have need for CBRN "plug and play" capability to allow interoperability and re-configurability across the enterprise. The requirement for net-centric, composable solutions provides the near term foundation for the Warfighter's ability to communicate his CBRN solutions and interoperate with other Service operational systems. It also supports a longer term ability to interoperate with related agencies and to reduce the Warfighter's CBRN footprint as technologies improve.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) JEM Operational Demonstrations and Exercises <i>FY 2009 Accomplishments:</i> Supported operational demonstrations and exercises. <i>FY 2010 Plans:</i> Continue to support operational demonstrations and exercises.	0.381	0.698	0.000	0.000	0.000
2) JEM Independent Verification, Validation, and Accreditation <i>FY 2009 Accomplishments:</i> Conducted independent verification, validation, and accreditation of JEM software and models. <i>FY 2010 Plans:</i> Conduct independent verification, validation, and accreditation of JEM software and models.	1.079	1.616	0.278	0.000	0.278

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2011 Base Plans:</i> Conduct independent verification, validation, and accreditation of JEM software and models.						
3) JEM Systems Engineering <i>FY 2009 Accomplishments:</i> Continued JEM Increment 1 Systems Engineering Tasks to include software updates, configuration management, human-system integration, security analysis and DoD architecture artifact development. Updated requirements and architecture analysis for supporting Science and Technology capabilities in preparation for JEM Increment 1. <i>FY 2010 Plans:</i> Continue to sustain JEM Increment 1 Systems Engineering Tasks to include software updates, configuration management, human-system integration, security analysis and DoD architecture artifact development.		1.121	0.794	0.000	0.000	0.000
4) JEM Program Management <i>FY 2009 Accomplishments:</i> Continued JEM program financial management, scheduling, planning and reporting. <i>FY 2010 Plans:</i> Continue JEM program financial management, scheduling, planning and reporting. <i>FY 2011 Base Plans:</i> Continue JEM program financial management, scheduling, planning and reporting.		1.913	1.947	0.233	0.000	0.233
5) JEM Program Development		2.309	2.229	0.478	0.000	0.478

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B. Accomplishments/Planned Program (\$ in Millions)								
				FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2010 Plans:</i> Analyze existing and future software architectures. Continue migrating JEM software to evolving host platforms (Service C2 systems). Incorporate Urban Dispersion Modeling enhancements, Missile Intercept, Backtracking to Source, enhanced STRATCOM Support, and Human Effects. Continue to review and evaluate existing JEM internal architecture for improved performance and potential operational cost savings.</p> <p><i>FY 2011 Base Plans:</i> Continue Science and Technology transition and improvement of JEM Increment 1 software. Analyze existing and future software architectures. Continue migrating JEM software to evolving host platforms (Service C2 systems). Incorporate Urban Dispersion Modeling enhancements, Missile Intercept, Backtracking to Source, enhanced STRATCOM Support, and Human Effects. Continue to review and evaluate existing JEM internal architecture for improved performance and potential operational cost savings.</p>								
9) JEM Developmental Test and Evaluation				1.091	2.073	0.439	0.000	0.439
<p><i>FY 2009 Accomplishments:</i> Continued to perform Governmental DT on JEM Increment 1 software. Verified that the JEM transitioned S&T code and model correctly and conducted test in support of follow-on accreditation and operational test. Completed interoperability, network and system security certifications of multiple service C4I/host systems and three computer operating systems (Windows XP, Vista and UNIX).</p> <p><i>FY 2010 Plans:</i> Continue to perform Governmental DT on JEM Increment 1 updates. Verify that the JEM transitioned S&T code and model correctly and conduct tests in support of follow-on accreditation and operational tests. Conduct interoperability, network and system security certifications of multiple service C4I/host systems and three computer operating systems (Windows XP, Vista and UNIX). Conduct verification and validation to ensure updates to evolving baseline don't affect JEM accreditation.</p>								

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B. Accomplishments/Planned Program (\$ in Millions)								
				FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
JOEF Test and Evaluation <i>FY 2009 Accomplishments:</i> Developed and tested interoperability of JOEF software with required systems (Increment 1). Planned and conducted Developmental and Operational Testing (DT/OT).								
13) JOEF JOEF Integrated Logistics Support <i>FY 2009 Accomplishments:</i> Planned and provided Integrated Logistics Support, including training, to the JOEF system (Increment 1).				0.320	0.000	0.000	0.000	0.000
14) JOEF JOEF Validation and verification (Increment 1) <i>FY 2009 Accomplishments:</i> Planned and conducted software validation and verification (Increment 1).				0.300	0.000	0.000	0.000	0.000
15) JOEF JOEF Integration with JEM and JWARN <i>FY 2009 Accomplishments:</i> Continued the integration with JEM, JWARN and database management systems (Increment 1).				0.300	0.000	0.000	0.000	0.000
16) JWARN Increment 1 JWARN Program Development				5.660	2.249	7.261	0.000	7.261

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2009 Accomplishments:</i> Continued JWARN program financial management, scheduling, planning and reporting.</p> <p><i>FY 2010 Plans:</i> Continue JWARN program financial management, scheduling, planning and reporting.</p> <p><i>FY 2011 Base Plans:</i> Continue JWARN program financial management, scheduling, planning and reporting.</p>						
<p>18) JWARN Increment 1 JWARN Operational demonstrations and tests.</p> <p><i>FY 2009 Accomplishments:</i> Conducted operational demonstrations and tests in support of Milestone "C" decision. Generated Multi-Service Operational Test and Evaluation (MOT&E) test results and reports.</p> <p><i>FY 2010 Plans:</i> Plan for, conduct and support operational demonstrations and tests in support of Follow-on Test and Evaluation (FOT&E). Generate test results and reports.</p> <p><i>FY 2011 Base Plans:</i> Prepare for and conduct operational demonstrations and tests.</p>		1.634	0.050	0.050	0.000	0.050
<p>19) JWARN Functional Qualification Tests (FQT)</p>		4.405	0.413	0.000	0.000	0.000

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2009 Accomplishments:</i> Conducted Increment 1 FQT to include a High Altitude Electro-Magnetic Pulse (HAEMP), MILSTD 810F environmental, and decontamination testing. Generated FQT results and reports.</p> <p><i>FY 2010 Plans:</i> Prepare for Increment 2 FQT.</p>						
<p>20) JWARN Increment 1 JWARN Total Package Fielding</p> <p><i>FY 2009 Accomplishments:</i> Developed and tested software to support JWARN Total Package Fielding.</p> <p><i>FY 2010 Plans:</i> Continue development and test of software to support JWARN Total Package Fielding.</p> <p><i>FY 2011 Base Plans:</i> Continue development and test of software to support JWARN Total Package Fielding.</p>		0.853	0.550	0.700	0.000	0.700
<p>21) JWARN Increment 1 JWARN Hardware Development</p> <p><i>FY 2009 Accomplishments:</i> Developed the wireless JWARN Component Interface Device (JCID) as required by the Services' Urgent Needs Statement (UNS). Produced 70 Engineering Development Models of the JCID.</p>		2.553	0.000	0.000	0.000	0.000
<p>22) SSA Policies, Standards and Guidelines</p>		0.422	0.233	0.094	0.000	0.094

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT IS5: <i>INFORMATION SYSTEMS (SDD)</i>
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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
32) SSA CBRN Interface Standards <i>FY 2009 Accomplishments:</i> Developed and maintained Common CBRN Interface standards, including a Common CBRN Sensor Interface (CCSI). <i>FY 2010 Plans:</i> Continue to maintain Common CBRN Interface standards, including a CCSI. <i>FY 2011 Base Plans:</i> Continue to maintain Common CBRN Interface standards, including a CCSI. Develop new interfaces as required.	0.624	0.402	0.162	0.000	0.162
33) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.	0.000	0.352	0.000	0.000	0.000
Accomplishments/Planned Programs Subtotals	45.694	27.301	13.844	0.000	13.844

C. Other Program Funding Summary (\$ in Millions)

Line Item	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	FY 2012	FY 2013	FY 2014	FY 2015	Cost To Complete	Total Cost
• G47101: <i>JOINT WARNING & REPORTING NETWORK (JWARN)</i>	4.375	6.551	6.903		6.903	8.078	5.590	8.183	8.423	Continuing	Continuing
• JC0208: <i>JOINT EFFECTS MODEL (JEM)</i>	5.546	3.482	3.482		3.482	0.000	0.000	3.369	3.568	Continuing	Continuing
	0.000	0.000	0.000		0.000	0.000	2.482	2.480	3.716	Continuing	Continuing

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C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011</u> <u>Base</u>	<u>FY 2011</u> <u>OCO</u>	<u>FY 2011</u> <u>Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To</u> <u>Complete</u>	<u>Total Cost</u>
• JC0209: <i>JOINT OPERATIONAL EFFECTS FEDERATION (JOEF)</i>											

D. Acquisition Strategy

JEM

The Joint Effects Model (JEM) is following an evolutionary acquisition approach that will allow rapid fielding of existing technologies while further research and development (R&D) continues in order to mature the technologies required for subsequent versions of JEM. It is now being fielded in increments of capabilities. Each increment will retain the functionality of the preceding increment. The JEM development effort will be aligned with the evolving JPED-CBD architectures and technologies, as well as with Service C2 systems. JEM is expected to develop three distinct increments of software. JEM will define and publish its web-services interface; the JEM interface will be the same on all systems, utilizing data definitions from the approved CBRN data model as appropriate. A cost plus award fee contract was awarded for the follow-on JEM contract for integration and development.

JOEF

Joint Operational Effects Federation (JOEF) is a planning tool to support deliberate and crisis planning. JOEF will be a near real-time course of action analysis tool developed in three increments. It will use a detailed CBRN hazard prediction model. Each block supports Aerial Ports of Debarkation (APODs), Sea Ports of Debarkation (SPODs), mobile forces, medical and automated Tactics, Techniques and Procedures (TTPs) in various levels of fidelity. Increment 1 will support deliberate planning for operational and strategic users in a Command, Control, Communications, Computers, Intelligence, Surveillance and Reconnaissance (C4ISR) common operating environment (COE)/Networked environment, Command and Control Personal Computers (C2PC), and crisis planning for the operational users in a COE/Networked environment. Increment 2 will support deliberate and crisis planning for the tactical users in COE/Networked, and Non-Networked environments; deliberate planning for operational and strategic users in a Non-Networked environment; and crisis planning for the operational users in a COE Networked and Non-Networked environments. Increment 2 also supports planning for consequence management and development of consequence management for military capabilities. Increment 3 will extend consequence management capabilities to include hot/allied nation military operations and civilian facilities.

JWARN

JWARN is being developed in two increments. JWARN Increment 1, will integrate JWARN capabilities into specified (Common Operating Environment (COE)-based) operational-level Service Command and Control (C2) systems at the Global Command and Control System (GCCS) stakeholder level, extend the integration effort

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<p>into the Service tactical (non COE-based) C2 systems, provide connectivity to legacy and newly developed sensors, and complete the development of JWARN in accordance with the threshold requirements contained in the Joint Requirements Oversight Council (JROC) approved Joint Operational Requirement Document (JORD) approved 5 July 2005, by JROC Memorandum 144-5 as refined by the JWARN Increment 1 Capabilities Production Document (CPD) approved 13 December 2007 by JROM 829-07. JWARN Increment 2 will extend these baseline capabilities to other C2 systems and new sensors as they are fielded; and further, will ensure CBRN warning and reporting capabilities remain synchronized with the changing demands of the Warfighter while keeping pace with evolving C2 systems.</p> <p>SSA</p> <p>The JPEO-CBD Software Support Activity (SSA) is a JPEO-CBD user support organization spanning and supporting all Joint Project Managers (JPMs) and JPEO-CBD Directorates. The SSA provides enterprise-wide services and coordination across all JPEO-CBD Programs of Record (PORs) that contain data or software, or are capable of linking to the Global Information Grid (GIG). The SSA facilitates interoperability, integration, and supportability of existing and developing IT and National Security Systems (NSS) across the JPEO and all JPMs.</p> <p>Phase 1a identifies JPEO-CBD JPMs and programs that deal with data or software, and have an IT component. This will be followed by coordination with the JPMs and programs to facilitate the concepts of interoperability, integration and supportability of enterprise-wide services. Next follows work with user communities to develop and demonstrate enterprise-wide common architectures, products and services. [BA5 - System Development and Demonstration].</p> <p>Phase 1b established management and control measures for tracking and reporting progress of the various elements described in Phases 1 and 2. This includes establishing, tracking, and performing configuration management of inventories and databases of IT systems and their states of interoperability and information assurance compliance. [BA6 - RDT&E Management Support].</p> <p>Phase 2 will support the application of the enterprise-wide architectures, products and services into the programs, with verification of compliance with the defined products and services. [BA7 - Operational Systems Development].</p> <p><u>E. Performance Metrics</u> N/A</p>		

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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Product Development (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** JEM - SW SB - JEM Hazard Prediction Model Development and Integration	C/CPAF	Northrop Grumman San Diego, CA	26.438	8.643	Jan 2010	0.000		0.000		0.000	0.000	35.081	0.000
** JWARN - SW S - JWARN System Development and Demonstration	C/CPAF	Northrop Grumman Winterpark, FL	12.238	2.000	Jan 2010	4.871	Jan 2011	0.000		4.871	0.000	19.109	0.000
** SSA - Product Development	MIPR	SPAWAR Systems Center San Diego, CA	4.641	1.020	Oct 2009	0.468	Oct 2010	0.000		0.468	0.000	6.129	0.000
Subtotal			43.317	11.663		5.339		0.000		5.339	0.000	60.319	0.000

Remarks

Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** JEM - ES S - IPT - System Engineering, Logistics and Program Support	MIPR	Various	15.203	1.998	Jan 2010	0.432	Jan 2011	0.000		0.432	0.000	17.633	0.000
** SSA - Support Costs	MIPR		4.821	1.444	Oct 2009	0.682	Oct 2010	0.000		0.682	0.000	6.947	0.000

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Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
		SPAWAR Systems Center San Diego, CA											
Subtotal			20.024	3.442		1.114		0.000		1.114	0.000	24.580	0.000

Remarks

Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** JEM - DTE SB - Hazard Prediction Model Development Test	MIPR	Various	6.035	2.073	Jan 2010	0.346	Jan 2011	0.000		0.346	0.000	8.454	0.000
OTE S - Hazard Prediction Model Developmental Test	MIPR	Various	5.555	2.219	Jan 2010	0.984	Jan 2011	0.000		0.984	0.000	8.758	0.000
OTHT SB - Hazard Prediction Model - IV&V	MIPR	Various	3.282	1.616	Jan 2010	0.000		0.000		0.000	0.000	4.898	0.000
** JWARN - OTHT SB - JWARN	MIPR	Various	31.247	0.161	Jan 2010	0.306	Jan 2011	0.000		0.306	0.000	31.714	0.000
** SSA - Test and Evaluation	MIPR	SPAWAR Systems Center	3.460	0.485	Oct 2009	0.148	Oct 2010	0.000		0.148	0.000	4.093	0.000

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Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
		San Diego, CA											
Subtotal			49.579	6.554		1.784		0.000		1.784	0.000	57.917	0.000

Remarks

Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** JEM - PM/MS S - Program Office - Planning and Programming	MIPR	SPAWAR Systems Command San Diego, CA	6.613	1.945	Oct 2009	0.233	Oct 2010	0.000		0.233	0.000	8.791	0.000
** JWARN - PM/MS S - JWARN Management Support	MIPR	Various	21.666	3.001	Jan 2010	5.328	Jan 2011	0.000		5.328	0.000	29.995	0.000
** SSA - Management Services	MIPR	SPAWAR Systems Center San Diego, CA	3.018	0.344	Oct 2009	0.046	Oct 2010	0.000		0.046	0.000	3.408	0.000
** ZSBIR - SBIR/STTR - Aggregated from ZSBIR-SBIR/STTR	MIPR	HQ AMC, Alexandria	0.000	0.352		0.000		0.000		0.000	0.000	0.352	0.000
Subtotal			31.297	5.642		5.607		0.000		5.607	0.000	42.546	0.000

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Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
Remarks													
			Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
Project Cost Totals			144.217	27.301		13.844		0.000		13.844	0.000	185.362	0.000

Remarks

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
** JEM - Increment 1 - Pre-planned Product Improvement (P3I)	■	■	■	■	■	■	■	■	■	■	■																	
Increment 1 - Production and Deployment	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■												
Increment 1 - Developmental Maintenance	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■												
Increment 1 - Follow-on Test and Evaluation				■	■	■																						
Increment 2 - Material Development Decision (MDD)					■																							
Increment 2 - Technology Development	■	■	■	■	■	■	■	■	■	■																		
Increment 2 - Analysis of Alternatives					■	■	■	■	■																			
Increment 2 - DT (Cont)							■	■	■	■	■	■	■	■	■	■	■	■										
Increment 2 - DT (Gov't)							■	■	■	■	■	■	■	■	■	■	■	■	■	■								
Increment 2 - Engineering and Manufacturing Development									■	■	■	■	■															
Increment 2 - Milestone B (MS B)															■													
Increment 2 - Milestone C (MS C)																			■									
Increment 2 - Multi-Service Operational Test and Evaluation (MOT&E)/LOG Demo																				■								
Increment 2 - Standalone Full Rate Production (FRP)																							■					
** JOEF - Increment 1 - Tech Reviews	■	■																										
Increment 1 - DT							■																					
Increment 1 - Operational Assessment																											■	

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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015							
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4				
Provide Enterprise Architecture Products and Services	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Provide Information Assurance Site Compliance Testing	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Provide Integration and Test, M&S, VV&A Certification and Accreditation	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Demonstrate Technology Transition Capabilities	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Provide CM Services for Common User Products and Services	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Provide Net-Centric Assessment and assist programs with implementation of policy	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Develop and provide CBRN Data Model implementation guidance, including reference implementations	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Architecture advisory services to support Warfighter Enterprise and Program Integrated Architectures	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Demonstrate, Verify, Test Technology Transition capabilities especially for Common Components and Services	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Provide Information Assurance Certification/Acceptance products/services, including compliance testing	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015							
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4				
Provide Modeling, Simulation, VV&A, Integration/Test support and interoperability demonstrations.	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Provide FISMA and J6 Interoperability certification support	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Provide CBRN Interface Standards, including reference implementations, e.g. Common CBRN Sensor Interface	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Sustain CBRN Data Model	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Sustain CCSI, including investigation, as an industry standard	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■

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Schedule Details

Event	Start		End	
	Quarter	Year	Quarter	Year
** JEM - Increment 1 - Pre-planned Product Improvement (P3I)	3	2008	3	2011
Increment 1 - Production and Deployment	4	2007	4	2012
Increment 1 - Developmental Maintenance	3	2008	4	2012
Increment 1 - Follow-on Test and Evaluation	4	2009	2	2010
Increment 2 - Material Development Decision (MDD)	1	2010	1	2010
Increment 2 - Technology Development	3	2008	2	2011
Increment 2 - Analysis of Alternatives	1	2010	1	2011
Increment 2 - DT (Cont)	3	2010	2	2013
Increment 2 - DT (Gov't)	3	2010	3	2013
Increment 2 - Engineering and Manufacturing Development	1	2011	2	2012
Increment 2 - Milestone B (MS B)	2	2012	2	2012
Increment 2 - Milestone C (MS C)	2	2013	2	2013
Increment 2 - Multi-Service Operational Test and Evaluation (MOT&E)/LOG Demo	4	2013	4	2013
Increment 2 - Standalone Full Rate Production (FRP)	2	2014	2	2014
** JOEF - Increment 1 - Tech Reviews	2	2006	2	2009
Increment 1 - DT	3	2010	3	2010
Increment 1 - Operational Assessment	3	2013	3	2013
Increment 1 - Multi-Service Operational Test & Evaluation (MOTE)	2	2014	2	2014

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Event	Start		End	
	Quarter	Year	Quarter	Year
Increment 1 - Milestone C (Limited Deployment)	2	2014	2	2014
Increment 1 - Initial Operational Capability (IOC)	1	2015	1	2015
Increment 1 - Full Operational Capability (FOC)	4	2015	4	2015
** JWARN - JWARN Inc 1 - First Article Test	4	2008	1	2009
JWARN Inc 1 - Multi-Service Operational Test & Evaluation (Software)	4	2008	2	2009
JWARN Inc 1 - Initial Operational Capability (Software)	1	2010	3	2010
JWARN Inc 1 - Full Rate Production Milestone Decision	2	2010	2	2010
JWARN Inc 1 - Full Rate Production	4	2010	2	2013
JWARN Inc 1 - Full Operational Capability	2	2011	2	2011
JWARN Inc 1 - Initial Operational Test and Evaluation (Hardware)	4	2010	4	2010
JWARN Inc 1 - Initial Operational Capability (Hardware)	1	2011	4	2011
** SSA - Provide Data Model Implementation Guidance	1	2008	4	2015
Provide Enterprise Architecture Products and Services	3	2007	4	2015
Provide Information Assurance Site Compliance Testing	3	2006	4	2015
Provide Integration and Test, M&S, VV&A Certification and Accreditation	2	2007	4	2015
Demonstrate Technology Transition Capabilities	1	2008	4	2015
Provide CM Services for Common User Products and Services	1	2008	4	2015
Provide Net-Centric Assessment and assist programs with implementation of policy	4	2007	4	2015
Develop and provide CBRN Data Model implementation guidance, including reference implementations	1	2008	4	2015
	1	2008	4	2015

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT IS5: <i>INFORMATION SYSTEMS (SDD)</i>
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Event	Start		End	
	Quarter	Year	Quarter	Year
Architecture advisory services to support Warfighter Enterprise and Program Integrated Architectures				
Demonstrate, Verify, Test Technology Transition capabilities especially for Common Components and Services	1	2008	4	2015
Provide Information Assurance Certification/Acceptance products/services, including compliance testing	1	2008	4	2015
Provide Modeling, Simulation, VV&A, Integration/Test support and interoperability demonstrations.	1	2008	4	2015
Provide FISMA and J6 Interoperability certification support	1	2008	4	2015
Provide CBRN Interface Standards, including reference implementations, e.g. Common CBRN Sensor Interface	1	2008	4	2015
Sustain CBRN Data Model	1	2008	4	2015
Sustain CCSI, including investigation, as an industry standard	1	2008	4	2015

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program									DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>				R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>				PROJECT MB5: <i>MEDICAL BIOLOGICAL DEFENSE (SDD)</i>			
COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
MB5: <i>MEDICAL BIOLOGICAL DEFENSE (SDD)</i>	87.676	57.558	141.680	0.000	141.680	161.732	159.144	141.481	111.671	Continuing	Continuing
Quantity of RDT&E Articles											

A. Mission Description and Budget Item Justification

This project (MB5) contains Engineering and Manufacturing Development (EMD) on efforts (post Milestone B), which provide a rapid response capability from identification of pathogens to the delivery of medical countermeasures. Specifically, this project includes: the Critical Reagents Program (CRP), the Joint Vaccine Acquisition Program (JVAP) which includes vaccines for Recombinant Botulinum A/B and Plague, and The Transformational Medical Technology Initiative (TMTI) program.

The Critical Reagents Program's (CRP) strategy establishes a core research and development capability to develop biological threat agent, genomic reference materials (antigens, nucleic acids, and antibodies) and detection and diagnostic assays for biothreat agent detection that shall be horizontally inserted across multiple detection and diagnostic platforms. In addition, this strategy will implement a formal, validated advanced development process to transition new assays into production and integration with the appropriate detection/diagnostic platform.

The Transformational Medical Technologies Initiative (TMTI) was launched to respond to the threat of emerging or intentionally bioengineered biological threats. TMTI's mission is to protect the Warfighter from genetically engineered biological threats by providing a rapid response capability from identification of pathogens to the delivery of medical countermeasures. This mission is accomplished by developing broad spectrum (multi-agent) therapeutics against biological warfare (BW) agents (e.g, one drug that treats multiple agents). The development of broad spectrum therapeutics involves developing a capability to treat exposure to hemorrhagic fever viruses (HFVs) (e.g. Ebola virus) and intracellular bacterial pathogens (ICBs) (e.g. Tularemia). Efforts are further classified as host-directed therapeutics (e.g, drugs that target common pathways within a human to prevent or treat a variety of diseases) or pathogen-directed therapeutics (e.g., drugs that attack a common pathway found in multiple threat agents). Attrition is high throughout the drug development process. Less than 10% of all preclinical compounds become a licensed drug. Causes for attrition include scientific failures, Food and Drug Administration (FDA) rejection at major milestone reviews, and loss through down-selection at DoD Milestone Decision points. The development of medical countermeasures is an arduous process that requires extensive interaction with the FDA, from pre-clinical research to safety tests in human subjects (Phase I clinical studies), efficacy tests in humans/animals (Phase II clinical studies or pivotal animal efficacy studies), and expanded safety or efficacy studies (Phase III clinical studies), which culminate with a request to the FDA to license, market, and produce a drug. This interaction between the Department of Defense (DoD) and the FDA results in a coordinated, unified, and safe effort.

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT MB5: <i>MEDICAL BIOLOGICAL DEFENSE (SDD)</i>
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The Joint Vaccine Acquisition Program (JVAP) under Chemical Biological Medical Systems (CBMS) funds the EMD phase of vaccines that are directed against validated biological warfare (BW) weapons to include bacteria, viruses, and toxins of biological origin. Effective medical countermeasures to negate the threat of these BW agents are urgently needed. Vaccines have been identified as the most efficient countermeasure against the validated threat of BW weapons. Efforts for medical biological defense product development involve production scale-up studies and validation, non-clinical studies, consistency manufacturing, and expanded clinical human safety studies. The results of these efforts, and those conducted during the EMD phase, will be used to submit a Biologic License Application (BLA) to the Food and Drug Administration (FDA) for product licensure. To evaluate vaccine effectiveness, pivotal animal studies will be conducted concurrently with the Phase 3 clinical trial to satisfy the requirements of the FDA's "Animal Rule". Upon FDA licensure, the product will transition to full-scale licensed production. Products under development in this budget item include Recombinant Botulinum A/B and Plague vaccines.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) CRP <i>FY 2009 Accomplishments:</i> Developed biological threat agent reference materials (new strains of Yersinia and Burkholderia). <i>FY 2010 Plans:</i> Continue development/expansion of biological select agents reference materials to known and emerging threats. <i>FY 2011 Base Plans:</i> Continue development/expansion of biological select agents reference materials to known and emerging threats.	2.217	1.158	1.278	0.000	1.278
2) CRP <i>FY 2009 Accomplishments:</i> Integrated new assays into CRP Catalog (PCR FastBlock assays, Positive Plasmid Controls and ECL System Reagents).	1.127	0.679	0.733	0.000	0.733

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT MB5: <i>MEDICAL BIOLOGICAL DEFENSE (SDD)</i>				
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2010 Plans:</i> Continue development of immunoassays and nucleic acid based genomic assays to support fielded and developmental systems.</p> <p><i>FY 2011 Base Plans:</i> Continue development of immunoassays and nucleic acid based genomic assays to support fielded and developmental systems.</p>						
<p>3) CRP</p> <p><i>FY 2009 Accomplishments:</i> Completed implementation of formal Quality Assurance/Quality Control(QA/QC) program to encompass systems engineering, validation, Development Testing (DT), and Operational Testing (OT) for the transition of fielding of biological detection assays.</p> <p><i>FY 2010 Plans:</i> Continue QA/QC testing to encompass the transition and fielding of biological detection assays.</p> <p><i>FY 2011 Base Plans:</i> Continue QA/QC testing to encompass the transition and fielding of biological detection assays.</p>		3.568	2.206	2.358	0.000	2.358
<p>4) CRP</p> <p><i>FY 2009 Accomplishments:</i> Continued development of guidelines to achieve International Organization for Standardization (ISO) for select biological threat agent reference materials.</p> <p><i>FY 2010 Plans:</i> Finalize implementation plan to achieve ISO certification.</p>		0.523	0.312	0.337	0.000	0.337

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2011 Base Plans:</i> Achieve and maintain ISO certification.						
5) TMTI Broad Spectrum Medical Countermeasures: Activities during this phase will include Phase 2 clinical trials, which will test the efficacy of therapeutics. Because the therapeutics sought are for biological warfare (BW) indications, efficacy testing on humans cannot be performed and will be tested in animals only. Because of the lack of human efficacy data, trials in animals are considered to be "Pivotal Animal Efficacy Studies" and are most commonly performed on at least two species of animals, to include non-human primates. Activities will also include expanded human and animal safety and/or animal efficacy studies as directed by the Food and Drug Administration (FDA) to support a New Drug Application (NDA) submission. <i>FY 2011 Base Plans:</i> Initiate pivotal animal efficacy studies and any expanded safety or efficacy studies as directed by the FDA for up to three candidate drugs following review and approval of new drug candidates at Milestone B. Critical activities will include dose/schedule and administration, assay validation, and efficacy studies in animals to demonstrate a favorable impact on clinical endpoints. Final formulation and dose will be determined by these studies. Pending discussions with the FDA, additional data may be required beyond that collected in initial Phase I clinical studies and pivotal animal efficacy studies to satisfy safety and/or efficacy requirements for NDA submission and licensure.		0.000	0.000	48.419	0.000	48.419
6) TMTI Platform Technologies: Exercises will commence on the platform technologies as an integrated system, and the bioinformatics system, developed with Science and Technology (S&T) funding, to evaluate and determine their capacity to support TMTI's capability goals. Platform technologies will continue to be integrated. Platforms will be exercised and evaluated as a system.		0.000	0.000	17.229	0.000	17.229

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2011 Base Plans:</i> Plan and execute up to two exercises and evaluations. Data will be collected into lessons learned and analyzed with the goal of improving the integration of the platforms. Analysis will be performed to develop a timeline for the response capability. Continue to evaluate the bioinformatics system for overall architecture, connectivity, processing capability, and user friendliness. Analyze lessons learned from each exercise and incorporate them into future exercises in order to improve countermeasure efficacy and shorten the time required to produce an approved countermeasure for an unknown or genetically modified pathogen.</p>						
<p>7) JVAP - Recombinant Botulinum Vaccine <i>FY 2009 Accomplishments:</i> Completed execution of Phase 1b clinical trial.</p>		0.874	0.000	0.000	0.000	0.000
<p>8) JVAP - Recombinant Botulinum Vaccine <i>FY 2009 Accomplishments:</i> Continued manufacturing process validation and validation of formulation, filled and finished process for serotypes A and B. <i>FY 2010 Plans:</i> Continue manufacturing process validation and validation of formulation, fill and finish process for serotypes A and B. <i>FY 2011 Base Plans:</i> Continue manufacturing process validation and validation of formulation, fill and finish process for serotypes A and B.</p>		18.059	22.874	28.668	0.000	28.668
<p>9) JVAP - Recombinant Botulinum Vaccine</p>		1.053	2.788	5.323	0.000	5.323

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2009 Accomplishments:</i> Continued non-clinical testing. Continued requirement for safeguarding biological select agents and toxins mandated by DoD Directive 5210.88.</p> <p><i>FY 2010 Plans:</i> Continue non-clinical testing. Continue requirement for safeguarding biological select agents and toxins mandated by DoD Directive 5210.88.</p> <p><i>FY 2011 Base Plans:</i> Continue non-clinical testing. Continue requirement for safeguarding biological select agents and toxins mandated by DoD Directive 5210.88.</p>						
10) JVAP - Recombinant Botulinum Vaccine		9.378	4.979	2.139	0.000	2.139
<p><i>FY 2009 Accomplishments:</i> Continued Phase 2 clinical trial.</p> <p><i>FY 2010 Plans:</i> Continue Phase 2 clinical trial.</p> <p><i>FY 2011 Base Plans:</i> Continue Phase 2 clinical trial.</p>						
11) JVAP - Plague Vaccine		12.956	1.453	0.000	0.000	0.000
<p><i>FY 2009 Accomplishments:</i> Continued large scale manufacturing process development.</p> <p><i>FY 2010 Plans:</i> Continue and complete large scale manufacturing process development.</p>						

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2010 Plans:</i> Continue large scale manufacturing process validation. <i>FY 2011 Base Plans:</i> Continue large scale manufacturing process validation.						
15) JVAP - Plague Vaccine <i>FY 2009 Accomplishments:</i> Provided strategic/tactical planning, government systems engineering, program/financial management, costing, technology assessment, contacting, scheduling, acquisition oversight and technical support. <i>FY 2010 Plans:</i> Provide strategic/tactical planning, government systems engineering, program/financial management, costing, technology assessment, contacting, scheduling, acquisition oversight and technical support. <i>FY 2011 Base Plans:</i> Provide strategic/tactical planning, government systems engineering, program/financial management, costing, technology assessment, contacting, scheduling, acquisition oversight and technical support.		4.484	2.888	4.298	0.000	4.298
16) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.		0.000	0.746	0.000	0.000	0.000
Accomplishments/Planned Programs Subtotals		87.676	57.558	141.680	0.000	141.680

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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C. Other Program Funding Summary (\$ in Millions)

Line Item	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	FY 2012	FY 2013	FY 2014	FY 2015	Cost To Complete	Total Cost
• JM0001: <i>JOINT BIO AGENT IDENT AND DIAG SYSTEM (JBAIDS)</i>	0.479	0.000	5.571		5.571	0.000	0.000	0.000	0.000	Continuing	Continuing
• JX0005: <i>DOD BIOLOGICAL VACCINE PROCUREMENT</i>	38.109	12.701	12.824		12.824	3.385	3.466	56.416	98.759	Continuing	Continuing
• JX0210: <i>CRITICAL REAGENTS PROGRAM (CRP)</i>	0.000	0.000	0.994		0.994	0.993	0.993	0.992	0.000	Continuing	Continuing
• MB4: <i>MEDICAL BIOLOGICAL DEFENSE (ACD&P)</i>	7.910	102.437	136.975		136.975	130.718	131.347	115.985	113.566	Continuing	Continuing

D. Acquisition Strategy

CRP

The Critical Reagents Program's (CRP) strategy establishes a core research and development capability to develop biological threat agent, genomic reference materials (antigens, nucleic acids, and antibodies) and detection and diagnostic assays for biothreat agent detection that shall be horizontally inserted across multiple detection and diagnostic platforms. In addition, this strategy will implement a formal, validated advanced development process to transition new assays into production and integration with the appropriate detection/diagnostic platform.

TMTI

Transformational Medical Technology Initiative's (TMTI) ultimate goal is the delivery of FDA-licensed, therapeutics to the Warfighter. This goal can be reached through any one of the following three acquisition approaches: 1) through the discovery of new drugs; 2) through application of new drug indications (i.e., through a commercial off-the-shelf (COTS) approach); or, 3) through the re-engineering of previously developed drugs (i.e., through a Modified COTS approach). This may involve FDA-approved drugs or previously developed drug compounds that do not have an FDA license. Each of these approaches will require different entry points into both the drug development process and the defense acquisition management timeline. Moreover, each of these approaches will likely experience a different set of FDA regulatory requirements. In order to execute the overall acquisition strategy, TMTI has partnered with other elements within the DoD Chemical and Biological Defense Program, DoD agencies, private industry, and other DoD laboratories for the development of TMTI products. The contract types used to execute the program will depend on the circumstances, including maturity of the science, the legalities surrounding Intellectual Property (IP) and patent rights, and even the

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
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<p>size of the performer. Cost Plus Fixed Fee and Cost Plus Incentive Fee contracts will be used with traditional or nontraditional defense contractors for most advanced development contracts. Finally, developing platform technologies, such as modeling and simulation to predict drug-to-drug interaction effects prior to actual clinical trials, and the use of genetic sequencing and a bioinformatics backbone, are examples of how TMTI managers intend to augment private industry best practices to streamline the program management, test and evaluation, and overall TMTI product development.</p> <p>VAC BOT</p> <p>A prime systems contractor will function as the "responsible head" and license holder and will perform all ancillary, regulatory, quality assurance, and data management as required by the FDA. The current budget supports development through FDA licensure of a recombinant bivalent (A and B) botulinum vaccine. Other serotypes will be developed through an evolutionary approach, as funding becomes available.</p> <p>The management lead for the program shifted to Joint Vaccine Acquisition Program (JVAP) at Milestone A. The Advanced Component Development and Prototypes (ACD&P) phase included the manufacture of candidate current Good Manufacturing Practices (cGMP) lots, animal safety testing, and initial clinical trials. During this phase, the vaccine was evaluated for safety and immunogenicity in a small human trial (Phase 1).</p> <p>During the Engineering and Manufacturing Development (EMD) phase, the JVAP prime systems contract (PSC) will stabilize the vaccine formulation, validate the manufacturing processes and testing protocols, optimize the delivery systems and manufacture consistency lots. Phase 2 clinical trials are performed during this phase to provide additional safety data and determine dose and schedule. The Phase 3 clinical trial also is conducted during this phase to demonstrate safety in an expanded volunteer population. To evaluate efficacy, pivotal animal studies will be conducted concurrently with the Phase 3 clinical trial to satisfy FDA requirements for the "Animal Rule." The Milestone C, also the Low Rate Initial Production (LRIP) decision, will be conducted after the manufacturing process has been validated and consistency lots have been produced. At the Milestone C, approval is granted to produce the Initial Operational Capability (IOC) of vaccine material. A Biologics Licensure Application is submitted to the FDA with all clinical, nonclinical, and manufacturing data. The FDA grants licensure to products that are determined to be safe and efficacious.</p> <p>VAC PLG</p> <p>Chemical Biological Medical Systems (CBMS) was mitigating technical program risk in the Plague Vaccine program by temporarily supporting development of both a US vaccine candidate and a United Kingdom vaccine candidate. During the 2008 Resource Allocation Decision, the US Plague Vaccine candidate was selected for development through licensure under JVAP's Prime Systems Contract. A Project Arrangement is in place with the United Kingdom and Canada.</p>		

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
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<p>The management lead for the program shifted to JVAP at Milestone A. The Advanced Component Development and Prototypes (ACD&P) phase included the manufacture of candidate current Good Manufacturing Practices (cGMP) lots, animal safety testing, and initial clinical trials. During this phase, the vaccine was evaluated for safety and immunogenicity in a small human trial (Phase 1).</p> <p>During the Engineering and Manufacturing Development phase (EMD), the vaccine developer will stabilize the vaccine formulation, validate the manufacturing processes and testing protocols, optimize the delivery systems, and manufacture consistency lots. Phase 2 clinical trials are performed during this phase to provide additional safety data and determine dose and schedule. The Phase 3 clinical trial is also conducted during this phase to demonstrate safety in an expanded volunteer population. To evaluate efficacy, pivotal animal studies will be conducted concurrently with the Phase 3 clinical trial to satisfy the requirements of the FDA's "Animal Rule." The Milestone C, also the Low Rate Initial Production (LRIP) decision, will be conducted after the manufacturing process has been validated and consistency lots have been produced. At the Milestone C, approval is granted to produce the Initial Operational Capability (IOC) of vaccine material. A Biologics License Application is submitted to the FDA with all clinical, nonclinical, and manufacturing data. The FDA grants licensure to products that are determined to be safe and efficacious.</p> <p><u>E. Performance Metrics</u> N/A</p>		

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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Product Development (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** CRP - CRP - Scale-up of Select Biological Threat Agent Reference Materials	MIPR	USAMRIID Fort Detrick, MD & Dugway Proving Ground	8.101	0.511	Jan 2010	0.607	Jan 2011	0.000		0.607	0.000	9.219	0.000
CRP - Development of Select Biological Threat Agent Reference Materials and Assays	MIPR	RDECOM Edgewood, MD	1.538	0.151	Jan 2010	0.159	Jan 2011	0.000		0.159	0.000	1.848	0.000
** TMTI - HW C - Therapeutic development	C/CPIF	TBD Contract #1	0.000	0.000		8.493	Apr 2011	0.000		8.493	0.000	8.493	0.000
HW C - Therapeutic development	C/CPIF	TBD Contract #2	0.000	0.000		8.493	Apr 2011	0.000		8.493	0.000	8.493	0.000
SW S - Technologies	C/CPIF	TBD Technologies	0.000	0.000		6.044	Apr 2011	0.000		6.044	0.000	6.044	0.000
** VAC BOT - Manufacturing, Validation and Consistency Lot Production	C/CPAF	DynPort Vaccine Company Frederick, MD	16.175	12.470	Jan 2010	14.554	Jan 2011	0.000		14.554	0.000	43.199	0.000
** VAC PLG - Manufacturing, Validation, and Consistency Lot Production	C/CPAF	DynPort Vaccine Company Frederick, MD	66.552	5.261	Jan 2010	13.743	Jan 2011	0.000		13.743	0.000	85.556	0.000
Subtotal			92.366	18.393		52.093		0.000		52.093	0.000	162.852	0.000

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT MB5: <i>MEDICAL BIOLOGICAL DEFENSE (SDD)</i>
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Product Development (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			

Remarks
 RDECOM - Research, Development & Engineering Command
 USAMRIID - US Army Medical Research Institute of Infectious Disease
 NMRC - Naval Medical Research Center
 USAMRIID - US Army Medical Research Institute of Infectious Diseases
 DPG - Dugway Proving Ground

Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** CRP - CRP - Select Biological Threat Agent Reference Material Regulatory /Quality Assurance (QA) Support	MIPR	DTIC Edgewood, MD	0.556	0.090	Jan 2010	0.095	Jan 2011	0.000		0.095	0.000	0.741	0.000
CRP - Select Biological Threat Agent Reference Material Support	MIPR	USAMRIID Fort Detrick, MD; RDECOM	2.290	0.294	Jan 2010	0.309	Jan 2011	0.000		0.309	0.000	2.893	0.000
CRP - Select Biological Threat Agent Reference Material Regulatory/ Quality Assurance (QA) Support	MIPR	Dugway Proving Ground Dugway, UT	1.171	0.131	Jan 2010	0.138	Jan 2011	0.000		0.138	0.000	1.440	0.000
	C/CPIF	TBD Contract #1	0.000	0.000		6.066	Apr 2011	0.000		6.066	0.000	6.066	0.000

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT MB5: <i>MEDICAL BIOLOGICAL DEFENSE (SDD)</i>
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Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** TMTI - ES C - Regulatory Integration (Environmental and FDA Documentation) and Delivery System													
ES C - Regulatory Integration (Environmental and FDA Documentation) and Delivery System	C/CPIF	TBD Contract #2	0.000	0.000		6.071	Apr 2011	0.000		6.071	0.000	6.071	0.000
TD/D C - Technologies	C/CPIF	TBD Technologies	0.000	0.000		4.317	Apr 2011	0.000		4.317	0.000	4.317	0.000
** VAC BOT - Regulatory Integration (Environmental and FDA Documentation) and Delivery System	C/CPAF	DynPort Vaccine Company Frederick, MD	2.909	1.559	Jan 2010	1.819	Jan 2011	0.000		1.819	0.000	6.287	0.000
** VAC PLG - Regulatory Integration (Environmental and FDA Documentation) and Delivery System	C/CPAF	DynPort Vaccine Company Frederick, MD	11.631	1.155	Jan 2010	1.418	Jan 2011	0.000		1.418	0.000	14.204	0.000
Subtotal			18.557	3.229		20.233		0.000		20.233	0.000	42.019	0.000

Remarks
 DTIC - Defense Technical Information Center
 NMRC - Naval Medical Research Center
 RDECOM - Research, Development & Engineering Command

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
USAMRIID - US Army Medical Research Institute of Infectious Diseases DPG - Dugway Proving Ground													

Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** CRP - CRP - Conformance Testing of Select Biological Threat Agent Reference Materials and Assays	MIPR	Naval Medical Research Center Silver Spring, MD	2.333	0.162	Jan 2010	0.206	Jan 2011	0.000		0.206	0.000	2.701	0.000
CRP - Test & Evaluation of Select Biological Threat Agent Reference Materials and Assays	MIPR	USAMRIID Frederick, MD	3.143	0.222	Jan 2010	0.282	Jan 2011	0.000		0.282	0.000	3.647	0.000
CRP - Validation Program	C/CPFF	TBD	5.431	0.597	Apr 2010	0.740	Apr 2011	0.000		0.740	0.000	6.768	0.000
** TMTI - DTE C - Phase II and III Testing	C/CPIF	TBD Contract #1	0.000	0.000		9.706	Apr 2011	0.000		9.706	0.000	9.706	0.000
DTE C - Phase II and III Testing	C/CPIF	TBD Contract #2	0.000	0.000		9.706	Apr 2011	0.000		9.706	0.000	9.706	0.000
Technologies	C/CPIF	TBD Technologies	0.000	0.000		6.752	Apr 2011	0.000		6.752	0.000	6.752	0.000

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** VAC BOT - Testing, Evaluation, and Clinical Trials	C/CPAF	DynPort Vaccine Company Frederick, MD	14.883	10.378	Jan 2010	12.479	Jan 2011	0.000		12.479	0.000	37.740	0.000
** VAC PLG - Testing, Evaluation, and Clinical Trials	C/CPAF	DynPort Vaccine Company Frederick, MD	55.678	8.181	Jan 2010	11.174	Jan 2011	0.000		11.174	0.000	75.033	0.000
Subtotal			81.468	19.540		51.045		0.000		51.045	0.000	152.053	0.000

Remarks
 DTIC - Defense Technical Information Center
 NMRC - Naval Medical Research Center
 RDECOM - Research, Development & Engineering Command
 USAMRIID - US Army Medical Research Institute of Infectious Diseases

Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** CRP - Product Management Support	Allot	CBMS Frederick, MD	0.924	0.365	Oct 2009	0.541	Oct 2010	0.000		0.541	0.000	1.830	0.000
Product Management Support	SS/FFP	Goldbelt Raven LLC, Frederick	3.079	1.493	Jan 2010	1.444	Jan 2011	0.000		1.444	0.000	6.016	0.000
Chem Bio Medical Systems Office	Allot	CBMS Frederick, MD	2.285	0.249	Jul 2010	0.180	Jul 2011	0.000		0.180	0.000	2.714	0.000

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program										DATE: February 2010			
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Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
IT, Facility and Security Support	MIPR	RDECOM Edgewood, MD	0.194	0.090	Jan 2010	0.005	Jan 2011	0.000		0.005	0.000	0.289	0.000
** VAC BOT - PM/MS S - Joint Vaccine Acquisition Program Management	Allot	CBMS Frederick, MD	0.349	1.543	Jul 2010	2.738	Jul 2011	0.000		2.738	0.000	4.630	0.000
PM/MS S - Contractor Systems Engineering/ Program Management Support	SS/FFP	Goldbelt Raven LLC, Frederick	2.983	1.805	Apr 2010	1.163	Apr 2011	0.000		1.163	0.000	5.951	0.000
PM/MS S - Award Fee (Maximum 10.5%)	C/CPAF	DynPort Vaccine Company Frederick, MD	3.691	2.886	Jan 2010	3.377	Jan 2011	0.000		3.377	0.000	9.954	0.000
** VAC PLG - PM/MS S - Joint Vaccine Acquisition Program Management Office	Allot	CBMS Frederick, MD	4.977	0.960	Jan 2010	1.577	Jan 2011	0.000		1.577	0.000	7.514	0.000
PM/MS S - Program Management Support	Allot	JPEO Falls Church, VA	6.225	2.888	Jul 2010	3.534	Jul 2011	0.000		3.534	0.000	12.647	0.000
PM/MS S - Contractor Systems Engineering/ Program Management Support #2	SS/FFP	Goldbelt Raven LLC, Frederick	7.870	0.851	Apr 2010	1.576	Apr 2011	0.000		1.576	0.000	10.297	0.000
PM/MS S - Award Fee (Maximum 10.5%) #2	C/CPAF	DynPort Vaccine Company Frederick, MD	10.860	2.520	Jan 2010	2.174	Jan 2011	0.000		2.174	0.000	15.554	0.000
	MIPR	HQ	0.000	0.746		0.000		0.000		0.000	0.000	0.746	0.000

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT MB5: <i>MEDICAL BIOLOGICAL DEFENSE (SDD)</i>
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Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** ZSBIR - SBIR/STTR - Aggregated from ZSBIR-SBIR/STTR		AMC, Alexandria											
Subtotal			43.437	16.396		18.309		0.000		18.309	0.000	78.142	0.000

Remarks

	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
		Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
Project Cost Totals		235.828	57.558		141.680		0.000	141.680	0.000	435.066	0.000

Remarks

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT MB5: <i>MEDICAL BIOLOGICAL DEFENSE (SDD)</i>
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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
** CRP - CRP - Expand Select Biological Threat Agent Reference Materials	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■										
CRP - Development of ECL Immunoassays & PCR Genomic Assays	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■										
CRP - Development and Implementation of Quality Initiatives, Validation Program, and Systems Engineering	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■										
CRP - Implementation of ISO Guidelines into Select Biological Threat Agent Reference Materials	■	■	■	■	■	■	■	■	■	■	■	■																
** TMTI - Milestone B Decision (Hemorrhagic Fever Viruses)											■																	
Contract 1-2 Phase II Pivotal Animal Studies											■	■	■	■	■	■												
** VAC BOT - rBV A/B - Process Validation - Large Scale	■	■	■	■	■	■	■	■	■	■	■	■	■	■														
rBV A/B - Non-Clinical Testing	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■										
rBV A/B - Phase 1 Clinical Trial (A/B)	■																											
rBV A/B - Phase 2 Clinical Trial (A/B)	■	■	■	■	■	■	■	■	■	■	■	■	■	■														
rBV A/B - Consistency Lot Production													■	■	■	■	■											
rBV A/B - Milestone C/LRIP																					■							
rBV A/B - Phase 3 Clinical Trial (A/B)																					■	■	■	■	■	■	■	■
** VAC PLG - PLG - Non-Clinical Studies	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■				

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT MB5: <i>MEDICAL BIOLOGICAL DEFENSE (SDD)</i>
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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
PLG - Process Development - Large Scale	■	■	■	■	■	■																						
PLG - Phase 2 Clinical Trial	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■											
PLG - Process Validation - Large Scale	■	■	■	■	■	■	■	■	■	■	■	■	■															
PLG - Consistency Lot Production													■	■	■													
PLG - Milestone C/LRIP																■												
PLG - Phase 3 Clinical Trial																	■	■	■	■	■	■	■	■	■			
PLG - Biological Licensure Application (BLA) Submission																												■

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT MB5: <i>MEDICAL BIOLOGICAL DEFENSE (SDD)</i>
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Schedule Details

Event	Start		End	
	Quarter	Year	Quarter	Year
** CRP - CRP - Expand Select Biological Threat Agent Reference Materials	4	2003	2	2013
CRP - Development of ECL Immunoassays & PCR Genomic Assays	1	2003	2	2013
CRP - Development and Implementation of Quality Initiatives, Validation Program, and Systems Engineering	4	2006	2	2013
CRP - Implementation of ISO Guidelines into Select Biological Threat Agent Reference Materials	3	2007	4	2011
** TMTI - Milestone B Decision (Hemorrhagic Fever Viruses)	3	2011	3	2011
Contract 1-2 Phase II Pivotal Animal Studies	3	2011	4	2012
** VAC BOT - rBV A/B - Process Validation - Large Scale	1	2002	1	2012
rBV A/B - Non-Clinical Testing	2	2003	2	2013
rBV A/B - Phase 1 Clinical Trial (A/B)	3	2004	1	2009
rBV A/B - Phase 2 Clinical Trial (A/B)	4	2008	2	2012
rBV A/B - Consistency Lot Production	1	2012	1	2013
rBV A/B - Milestone C/LRIP	2	2013	2	2013
rBV A/B - Phase 3 Clinical Trial (A/B)	3	2013	1	2016
** VAC PLG - PLG - Non-Clinical Studies	2	2003	2	2014
PLG - Process Development - Large Scale	4	2004	2	2010
PLG - Phase 2 Clinical Trial	3	2006	1	2013
PLG - Process Validation - Large Scale	4	2007	1	2012

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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Event	Start		End	
	Quarter	Year	Quarter	Year
PLG - Consistency Lot Production	1	2012	3	2012
PLG - Milestone C/LRIP	4	2012	4	2012
PLG - Phase 3 Clinical Trial	1	2013	1	2015
PLG - Biological Licensure Application (BLA) Submission	1	2015	1	2015

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
MC5: <i>MEDICAL CHEMICAL DEFENSE (SDD)</i>	14.203	14.027	51.856	0.000	51.856	47.835	28.771	12.122	8.171	Continuing	Continuing
Quantity of RDT&E Articles											

A. Mission Description and Budget Item Justification

This project funds the development of medical materiel and other medical equipment items necessary to provide an effective capability for medical defense against chemical agent threats facing U.S. forces in the field. This project supports efforts in the Engineering and Manufacturing Development (EMD) phase of the acquisition strategy for prophylactic and therapeutic drugs, diagnostic equipment, and other life support equipment for protection against and management of chemical warfare agents. Project funds research and development of safety studies, manufacturing scale-up, process validation, drug interaction, performance test, and submission of the Food and Drug Administration (FDA) drug licensure application(s). This program currently funds: (1) Advanced Anticonvulsant System (AAS), which will be used as a treatment for seizures from exposure to nerve agents; (2) Bioscavenger Increment 2 (BSCAV Increment 2), which will be used as a prophylaxis against nerve agents; (3) Inhalational Atropine (IA), which will be used to treat continuing nerve agent-induced effects after the patient has been evacuated to a medical treatment facility; (4) Improved Nerve Agent Treatment System (INATS), which will be used as a treatment for nerve agent intoxication to include new indications for Pyridostigmine Bromide (PB) that will be integrated with current therapeutic regimens; and (5) Pharmaceutical Post Approval and Development Support (PPADS) - Soman Nerve Agent Pyridostigmine Pretreatment (SNAPP) used as a pretreatment against nerve agent poisoning. Time Temperature Indicators (TTI), Item Unique Identification (IUID), and Radio-Frequency Identification (RFID) will be part of the development effort for incorporation on all medical countermeasures being developed by CBMS-MITS. A TTI is a human readable tab that will provide the Warfighter immediate knowledge if the product is still useable or not. IUID and RFID labels placed on the product will improve inventory management and strategic purchasing, and enable reliable visibility, and capability-based operational readiness.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) AAS <i>FY 2009 Accomplishments:</i> Continued Phase 2 clinical safety studies.	3.491	0.176	0.000	0.000	0.000

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program			DATE: February 2010			
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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2011 Base Plans:</i> Initiate Phase 2 clinical safety studies.						
10) BSCAV <i>FY 2011 Base Plans:</i> Initiate BLA preparation and submission to Food and Drug Administration (FDA).		0.000	0.000	1.050	0.000	1.050
11) BSCAV NTA <i>FY 2011 Base Plans:</i> Initiate GLP animal studies to demonstrate efficacy against a broad spectrum of nerve agents including non-traditional agents.		0.000	0.000	8.878	0.000	8.878
12) BSCAV BSCAV <i>FY 2010 Plans:</i> Initiate proof of concept studies for alternative manufacturing technology. <i>FY 2011 Base Plans:</i> Continue proof of concept studies for alternative manufacturing technology.		0.000	0.800	1.005	0.000	1.005
13) IA <i>FY 2011 Base Plans:</i> Continue process development and current Good Manufacturing Practices (cGMP) requirements.		0.000	0.000	1.605	0.000	1.605
14) Inhalational Atropine (IA)		0.000	0.000	0.900	0.000	0.900

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2009 Accomplishments:</i> Provided strategic/tactical planning, government systems engineering, program/financial management, costing, technology, assessment, contracting, scheduling, acquisition oversight and technology support.						
20) INATS <i>FY 2011 Base Plans:</i> Initiate New Drug Application (NDA) preparation.		0.000	0.000	0.600	0.000	0.600
21) INATS INATS - Test candidate oxime against non-traditional agents. <i>FY 2011 Base Plans:</i> INATS - Test candidate oxime against non-traditional agents.		0.000	0.000	10.496	0.000	10.496
22) PPADS <i>FY 2010 Plans:</i> Develop a Time Temperature Indicator (TTI) capability for Soman Nerve Agent Pre-Treatment Pyridostigmine to provide visual indicator of product reliability.		0.000	0.738	0.000	0.000	0.000
23) PPADS <i>FY 2009 Accomplishments:</i> Initiated development of a cGMP process for the production of Sodium Thiosulfate and Sodium Nitrite as part of the cyanide antidote kit.		0.340	0.000	0.000	0.000	0.000
24) SBIR		0.000	0.180	0.000	0.000	0.000

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2010 Plans:</i> Small Business Innovative Research.					
Accomplishments/Planned Programs Subtotals	14.203	14.027	51.856	0.000	51.856

C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011 Base</u>	<u>FY 2011 OCO</u>	<u>FY 2011 Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To Complete</u>	<u>Total Cost</u>
• JM6500: <i>INHALATIONAL ATROPINE (IA)</i>	0.000	0.000	0.000		0.000	0.000	0.000	0.496	0.991	Continuing	Continuing
• JM6555: <i>IMPROVED NERVE AGENT TREATMENT SYSTEM (INATS)</i>	0.000	0.000	0.000		0.000	0.000	0.000	3.966	4.954	Continuing	Continuing
• JM6677: <i>ADVANCED ANTICONVULSANT SYSTEM (AAS)</i>	0.000	0.000	0.000		0.000	0.000	4.466	9.126	5.187	Continuing	Continuing

D. Acquisition Strategy

AAS

The Advanced Anticonvulsant System (AAS) will consist of the drug midazolam in an autoinjector. Midazolam, injected intramuscularly, will treat against seizures and prevent subsequent neurological damage caused by exposure to nerve agents. Midazolam is more water-soluble than diazepam (the currently fielded medication to control nerve agent-induced seizures) and terminates nerve agent-induced seizures more quickly than diazepam. AAS will not eliminate the need for other protective and therapeutic systems. AAS will be a replacement for the currently-fielded Convulsant Antidote, Nerve Agent (CANA) autoinjector, which uses diazepam.

BSCAV

The Bioscavenger acquisition strategy consists of a developmental program with three distinct increments.

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<p>Increment 1 is butyrylcholinesterase purified from human plasma (i.e., plasma-derived Bioscavenger or pBioscavenger). The Medical Identification and Treatment Systems (MITS) Joint Product Management Office exercises management oversight, and a commercial partner serves as the system integrator during the Technology Development Phase, which includes small scale manufacturing, pre-clinical animal studies, Investigational New Drug (IND) application, and Phase 1 human clinical safety studies.</p> <p>The Bioscavenger Increment 2 strategy includes a proof-of-concept study followed by an initial down-selection between two different technologies: Recombinant human butyrylcholinesterase (rHuBChE) and small synthetic molecule, awarded to two different contractors. The chosen technology, rHuBChE, will continue to a formal down-selection with the plasma-derived Bioscavenger at Milestone B prior to transition to the Engineering and Manufacturing Development (EMD) phase. Following Milestone B into EMD, MITS will continue to exercise management oversight with system integration support of a commercial partner to ensure manufacturing of the product is in accordance with Food and Drug Administration (FDA) regulations and guidelines. Prior to FDA licensure, the commercial partner will perform a Phase 2 human clinical safety study, definitive animal efficacy studies, and toxicology studies. The SDD phase will culminate in obtaining FDA licensure of the Bioscavenger. During the Production and Deployment phase, the MITS JPMO, in conjunction with a commercial partner, will pursue full rate and stockpile production and conduct any FDA-mandated post-marketing surveillance.</p> <p>Unlike Bioscavenger Increment 1 and 2 technology, where the Bioscavenger becomes ineffective after binding with nerve agents, Increment 3 will include products that continuously degrade nerve agents while retaining their effectiveness (catalytic Bioscavenger). Because the technology for Increment 3 is immature, a candidate product will not be ready for transition to advanced development until late in the next decade. Therefore, CBMS MITS is exploring alternative technologies to reduce the costs of producing Bioscavenger Increment 2.</p> <p>IA</p> <p>Medical Identification and Treatment Systems (MITS) Joint Product Management Office (JPMO) will manage the development of Inhalational Atropine (IA) for the DoD. For this Advanced Development effort, the competitively selected contractor will serve as the systems integrator throughout development and shall be responsible for conducting the activities associated with drug development in a manner consistent with eventual approval by the Food and Drug Administration (FDA), including: human clinical safety studies; pharmacokinetic studies; and validated manufacturing. The contractor shall sponsor the drug product to the FDA and hold all approvals and/or licenses.</p> <p>INATS</p> <p>Medical Identification and Treatment Systems (MITS) Joint Product Management Office will serve as the system integrator during the Technology Development Phase that includes pre-clinical animal studies and Phase 1 human clinical safety studies. After Milestone B, during the System Development and Demonstration Phase,</p>		

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0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	MC5: <i>MEDICAL CHEMICAL DEFENSE (SDD)</i>

MITS and/or a commercial partner (product dependent) will serve as the system integrator to ensure that products are manufactured in accordance with Food and Drug Administration (FDA) regulations and guidelines, appropriate Phase 2 human clinical safety and definitive animal efficacy studies are conducted, and required toxicology studies are performed. During the Production and Deployment Phase, FDA approval will be obtained and full rate and stockpile production will be pursued. Any FDA mandated post-marketing surveillance will be conducted.

E. Performance Metrics

N/A

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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Product Development (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** AAS - AAS - cGMP Manufacturing Requirements	C/CPIF	Meridian Medical Technologies Columbia, MD	11.018	1.409	Jan 2010	1.029	Jan 2011	0.000		1.029	0.000	13.456	0.000
** BSCAV - BSCAV Inc 2 - cGMP Manufacturing	C/CPIF	TBD	0.000	2.714	Apr 2010	6.010	Jan 2011	0.000		6.010	0.000	8.724	0.000
** IA - cGMP Manufacturing requirements	C/CPIF	TBD	0.000	0.000		0.945	Jan 2011	0.000		0.945	0.000	0.945	0.000
** INATS - INATS - cGMP Manufacturing	C/CPIF	TBD	0.000	1.597	Jan 2010	2.912	Jan 2011	0.000		2.912	0.000	4.509	0.000
INATS - NTA Study	C/CPIF	TBD	0.000	0.000		10.496	Jan 2011	0.000		10.496	0.000	10.496	0.000
Subtotal			11.018	5.720		21.392		0.000		21.392	0.000	38.130	0.000

Remarks

Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** AAS - AAS - Regulatory Integration and NDA Support Efforts	C/CPIF	Meridian Medical Technologies Columbia, MD	4.018	0.529	Jan 2010	0.391	Jan 2011	0.000		0.391	0.000	4.938	0.000
	C/CPIF	TBD	0.000	1.075	Apr 2010	2.626	Jan 2011	0.000		2.626	0.000	3.701	0.000

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Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** BSCAV - BSCAV Inc 2 - Regulatory Integration & Biologics License Application (BLA) Support Efforts													
** IA - Regulatory Integration and NDA support efforts	C/CPIF	TBD	0.000	0.000		0.390	Jan 2011	0.000		0.390	0.000	0.390	0.000
** INATS - INATS - Regulatory Integration and NDA Support Efforts	C/CPIF	TBD	0.000	0.543	Jan 2010	1.092	Jan 2011	0.000		1.092	0.000	1.635	0.000
Subtotal			4.018	2.147		4.499		0.000		4.499	0.000	10.664	0.000

Remarks

Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** AAS - AAS - GLP Animal Efficacy Studies	C/CPFF	Battelle Memorial Institute Columbus, OH	2.949	0.921	Jan 2010	0.698	Jan 2011	0.000		0.698	0.000	4.568	0.000
** BSCAV - BSCAV Inc 2 - Phase 2 Clinical	C/CPIF	TBD	0.000	2.034	Apr 2010	6.193	Jan 2011	0.000		6.193	0.000	8.227	0.000

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Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
Safety and GLP Animal Efficacy Studies													
BSCAV Inc 2 - Animal Efficacy Studies	C/CPIF	TBD	0.000	0.000		8.878	Jan 2011	0.000		8.878	0.000	8.878	0.000
** IA - Formulation and device development studies	C/CPIF	TBD	0.000	0.000		0.780	Jan 2011	0.000		0.780	0.000	0.780	0.000
** INATS - INATS - GLP Animal Efficacy & Phase 2 Clinical Safety Studies	C/CPIF	TBD	0.000	0.410	Jul 2010	3.963	Apr 2011	0.000		3.963	0.000	4.373	0.000
INATS - Large Scale Manufacturing	C/CPIF	TBD	0.000	0.000		0.881	Apr 2011	0.000		0.881	0.000	0.881	0.000
** PPADS - PPADS - Time Temperature Indicator (TTI) Capability	C/CPIF	TBD	0.000	0.738	Jan 2010	0.000		0.000		0.000	0.000	0.738	0.000
Subtotal			2.949	4.103		21.393		0.000		21.393	0.000	28.445	0.000

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Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** AAS - AAS - Product Management Support	MIPR	USAMMDA Fort Detrick, MD	0.689	0.165	Jan 2010	0.170	Jan 2011	0.000		0.170	0.000	1.024	0.000
AAS - Chem Bio Medical Systems	Allot	CBMS Frederick, MD	1.127	0.364	Apr 2010	0.221	Jan 2011	0.000		0.221	0.000	1.712	0.000
** BSCAV - BSCAV Inc 2 - Product Management Support	SS/FFP	Goldbelt Raven LLC, Frederick	0.000	0.502	Apr 2010	0.825	Jan 2011	0.000		0.825	0.000	1.327	0.000
BSCAV Inc 2 - Chem Bio Medical Systems	Allot	CBMS Frederick, MD	0.000	0.054	Apr 2010	0.081	Apr 2011	0.000		0.081	0.000	0.135	0.000
BSCAV Inc 2 - Joint Program Executive Office	Allot	JPEO Falls Church, VA	0.000	0.358	Apr 2010	0.542	Jan 2011	0.000		0.542	0.000	0.900	0.000
USAMMDA, Fort Detrick, MD	Allot	Fort Detrick MD	0.000	0.161	Apr 2010	0.178	Apr 2011	0.000		0.178	0.000	0.339	0.000
** IA - IA - Management Support	Allot	CBMS Frederick, MD	0.000	0.000		0.260	Jan 2011	0.000		0.260	0.000	0.260	0.000
IA - Management Support	Allot	JPEO Falls Church, VA	0.000	0.000		0.130	Jan 2011	0.000		0.130	0.000	0.130	0.000
** INATS - INATS - Product Management Support	SS/FFP	Goldbelt Raven LLC, Frederick	0.000	0.000		0.705	Jan 2011	0.000		0.705	0.000	0.705	0.000
INATS - Product Management Support	MIPR	USAMMDA Fort Detrick, MD	0.000	0.000		0.160	Apr 2011	0.000		0.160	0.000	0.160	0.000
INATS - Chem Bio Medical Systems	Allot	CBMS Frederick, MD	0.000	0.273	Jan 2010	0.300	Jan 2011	0.000		0.300	0.000	0.573	0.000
	Allot	JPEO	1.016	0.000		1.000	Jan 2011	0.000		1.000	0.000	2.016	0.000

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
INATS - Joint Program Executive Office		Falls Church, VA											
** ZSBIR - SBIR/STTR - Aggregated from ZSBIR-SBIR/STTR	MIPR	HQ AMC, Alexandria	0.000	0.180		0.000		0.000		0.000	0.000	0.180	0.000
Subtotal			2.832	2.057		4.572		0.000		4.572	0.000	9.461	0.000

Remarks

	Total Prior Years Cost	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
Project Cost Totals	20.817	14.027	51.856	0.000	51.856	0.000	86.700	0.000

Remarks

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
** AAS - AAS - Process development and cGMP Manufacturing Requirements	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■												
AAS - Formulation and Toxicology Studies	■	■																										
AAS - GLP Animal Efficacy Studies	■	■	■	■	■	■	■	■																				
AAS - Phase 2 Clinical Safety Studies	■	■	■	■	■	■																						
AAS - DT/OT for Packaging	■	■	■	■	■	■																						
AAS - New Drug Application (NDA) Preparation			■	■	■	■	■	■	■	■	■																	
AAS - MS C																■												
** BSCAV - BSCAV Inc. 2 - Large Scale Manufacturing, Process Development & Assay Validation	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■				
BSCAV Inc. 2 - Milestone B					■																							
BSCAV Inc. 2 - Conduct GLP Animal Efficacy Studies							■	■	■	■	■	■	■	■	■	■	■	■	■	■								
BSCAV Inc. 2 - Conduct Phase 2 Clinical Safety Studies									■	■	■	■	■	■	■	■	■	■	■	■	■	■						
BSCAV Inc. 2 - BLA Preparation and Submittal											■	■	■	■	■	■	■	■	■	■	■	■	■	■				
BSCAV -						■																						
** IA - IA - Milestone A					■																							
IA - Process Development and current Good Manufacturing Practices (cGMP) requirements						■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
IA - Formulation, analytical assay, and device development							■	■	■	■	■	■	■	■	■	■												
IA - Milestone B							■																					
IA - Phase 2 Clinical Safety studies													■	■	■	■	■	■	■	■								
IA - New Drug Application (NDA) Preparation and submission																	■	■	■	■								
IA - Process Development and cGMP requirements							■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
IA - Continue process development and cGMP requirements							■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
** INATS - INATS - Large Scale Manufacturing											■	■	■	■	■	■	■	■	■	■								
INATS - Phase 2 Clinical Safety Studies											■	■	■	■	■	■	■	■	■	■								
INATS - GLP Animal Efficacy Studies											■	■	■	■	■	■	■	■	■	■								
INATS - NDA Preparation and Submittal											■	■	■	■	■	■	■	■	■	■								
** PPADS - PPADS - Develop Time Temperature Indicator (TTI) Capability						■	■	■																				
PPADS - Cyanide Antidote Kit			■																									

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
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Schedule Details

Event	Start		End	
	Quarter	Year	Quarter	Year
** AAS - AAS - Process development and cGMP Manufacturing Requirements	4	2005	4	2012
AAS - Formulation and Toxicology Studies	3	2007	2	2009
AAS - GLP Animal Efficacy Studies	4	2007	3	2010
AAS - Phase 2 Clinical Safety Studies	3	2007	2	2010
AAS - DT/OT for Packaging	4	2008	2	2010
AAS - New Drug Application (NDA) Preparation	3	2009	3	2011
AAS - MS C	1	2013	1	2013
** BSCAV - BSCAV Inc. 2 - Large Scale Manufacturing, Process Development & Assay Validation	1	2008	4	2014
BSCAV Inc. 2 - Milestone B	2	2010	2	2010
BSCAV Inc. 2 - Conduct GLP Animal Efficacy Studies	3	2010	4	2013
BSCAV Inc. 2 - Conduct Phase 2 Clinical Safety Studies	1	2011	2	2014
BSCAV Inc. 2 - BLA Preparation and Submittal	4	2011	4	2014
BSCAV -	3	2010	3	2010
** IA - IA - Milestone A	2	2010	2	2010
IA - Process Development and current Good Manufacturing Practices (cGMP) requirements	3	2010	3	2015
IA - Formulation, analytical assay, and device development	3	2010	4	2012
IA - Milestone B	4	2010	4	2010

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Event	Start		End	
	Quarter	Year	Quarter	Year
IA - Phase 2 Clinical Safety studies	1	2012	1	2014
IA - New Drug Application (NDA) Preparation and submission	3	2013	2	2014
IA - Process Development and cGMP requirements	3	2010	3	2015
IA - Continue process development and cGMP requirements	3	2010	3	2015
** INATS - INATS - Large Scale Manufacturing	3	2011	1	2014
INATS - Phase 2 Clinical Safety Studies	3	2011	3	2013
INATS - GLP Animal Efficacy Studies	3	2011	3	2013
INATS - NDA Preparation and Submittal	4	2011	3	2013
** PPADS - PPADS - Develop Time Temperature Indicator (TTI) Capability	2	2010	4	2010
PPADS - Cyanide Antidote Kit	4	2009	4	2009

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0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>			PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>				MR5: <i>MEDICAL RADIOLOGICAL DEFENSE (SDD)</i>				
COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
MR5: <i>MEDICAL RADIOLOGICAL DEFENSE (SDD)</i>	3.002	8.276	1.143	0.000	1.143	4.817	2.265	0.000	0.000	Continuing	Continuing
Quantity of RDT&E Articles											

A. Mission Description and Budget Item Justification

This project funds the advanced development of candidate therapeutic medical countermeasures to mitigate the consequences of exposure to ionizing radiation due to nuclear or radiological attacks. Exposure to ionizing radiation causes damage to blood-forming cells (hematopoietic system) and gastrointestinal system, leading to Acute Radiation Syndrome (ARS). Medical countermeasures must be approved by the Food and Drug Administration (FDA) for human use prior to fielding. Testing the efficacy of candidate drugs against normally lethal radiation exposure cannot be conducted in humans; therefore, surrogate animal models must be used to obtain FDA approval.

Medical Radiation Countermeasures (MRADC) efforts include multiple countermeasures required to restore casualties to pre-exposure health and to protect U.S. Forces against injury caused by exposure to radiation. MRADC shall reverse or limit radiation injury resulting in increase survival, decreased incapacity, and sustained operational effectiveness. In addition, MRADC shall be effective against a broad range of radiation sources and types, and shall be useable in the battle space, including evacuation.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) MRADC MRADC <i>FY 2010 Plans:</i> Initiate non-clinical animal efficacy studies.	0.000	2.080	0.368	0.000	0.368

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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2011 Base Plans:</i> Continue non-clinical animal efficacy studies.						
2) MRADC <i>FY 2009 Accomplishments:</i> Provided strategic/tactical planning, government systems engineering, program/financial management, costing, technology assessment, contracting, scheduling, acquisition oversight and technical support.		3.002	0.000	0.000	0.000	0.000
3) MRADC <i>FY 2010 Plans:</i> Initiate process development and current Good Manufacturing Practices (cGMP) manufacturing requirements. <i>FY 2011 Base Plans:</i> Continue process development and current Good Manufacturing Practices (cGMP) manufacturing requirements.		0.000	3.583	0.395	0.000	0.395
4) MRADC <i>FY 2010 Plans:</i> Initiate product formulation, storage, and delivery system on 2 candidates. <i>FY 2011 Base Plans:</i> Continue product formulation, storage, and delivery system on 2 candidates.		0.000	2.505	0.380	0.000	0.380
5) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.		0.000	0.108	0.000	0.000	0.000

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT MR5: <i>MEDICAL RADIOLOGICAL DEFENSE (SDD)</i>
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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
Accomplishments/Planned Programs Subtotals	3.002	8.276	1.143	0.000	1.143

C. Other Program Funding Summary (\$ in Millions)

N/A

D. Acquisition Strategy

MRADC

Medical Identification and Treatment Systems (MITS) Joint Product Management Office will manage the development of Medical Radiation Countermeasures (MRADC) for the Department of Defense (DoD). A contractor will serve as the product integrator throughout development and shall be responsible for conducting activities associated with drug development in a manner consistent with eventual approval by the Food and Drug Administration (FDA). The contractor shall sponsor the drug to the FDA and hold all approvals and/or licenses. The Technology Development phase includes pre-clinical studies and Phase 1 human clinical safety studies. During the Engineering and Manufacturing Development (EMD) phase, large scale manufacturing, Phase 2 human clinical safety studies and definitive animal efficacy studies will be conducted. FDA approval of the countermeasure is an exit criterion for the EMD phase. During the Production and Deployment Phase, sufficient quantities of product to meet Initial Operational Capability and Full Operational Capability will be purchased. Subsequent purchases will be made by the Defense Logistics Agency. Any post-marketing surveillance requested by the FDA will be conducted.

MRADC will be developed using a system-of-systems approach to provide a full spectrum capability to protect against the radiation threat. Individual countermeasure solutions will be developed using a single step to a full capability (FDA approval) strategy. The DoD is working very closely with the Department of Health and Human Services (HHS), which also has a radiation countermeasure program. The establishment of an interagency working group provides oversight and guidance to both agency programs to ensure that their efforts are non-duplicative and are directed to meeting the requirement of both agencies.

E. Performance Metrics

N/A

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT MR5: <i>MEDICAL RADIOLOGICAL DEFENSE (SDD)</i>
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Product Development (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** MRADC - MRADC - cGMP Manufacturing	C/CPIF	TBD	0.000	2.346	Jan 2010	0.377	Jan 2011	0.000		0.377	0.000	2.723	0.000
MRADC - Product Formulation, Storage and Delivery System	C/CPIF	TBD	0.000	2.219	Jan 2010	0.200	Jan 2011	0.000		0.200	0.000	2.419	0.000
Subtotal			0.000	4.565		0.577		0.000		0.577	0.000	5.142	0.000

Remarks

Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** MRADC - MRADC - Regulatory Integration and NDA Support Efforts	C/CPIF	TBD	0.000	0.000		0.100	Jan 2011	0.000		0.100	0.000	0.100	0.000
Subtotal			0.000	0.000		0.100		0.000		0.100	0.000	0.100	0.000

Remarks

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT MR5: <i>MEDICAL RADIOLOGICAL DEFENSE (SDD)</i>
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Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** MRADC - MRADC - Definitive Animal Efficacy studies	C/CPIF	TBD	0.000	2.087	Jan 2010	0.200	Jan 2011	0.000		0.200	0.000	2.287	0.000
Subtotal			0.000	2.087		0.200		0.000		0.200	0.000	2.287	0.000

Remarks

Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** MRADC - MRADC - Product Management Support	SS/FFP	Goldbelt Raven LLC, Frederick	0.657	0.705	Jan 2010	0.000		0.000		0.000	0.000	1.362	0.000
MRADC - Chem Bio Medical Systems	Allot	CBMS Frederick, MD	0.000	0.659	Apr 2010	0.100	Apr 2011	0.000		0.100	0.000	0.759	0.000
MRADC - Product Management Services	MIPR	USAMMDA Ft Detrick, MD	0.145	0.152	Oct 2009	0.166	Oct 2010	0.000		0.166	0.000	0.463	0.000
** ZSBIR - SBIR/STTR - Aggregated from ZSBIR-SBIR/STTR	MIPR	HQ AMC, Alexandria	0.000	0.108		0.000		0.000		0.000	0.000	0.108	0.000
Subtotal			0.802	1.624		0.266		0.000		0.266	0.000	2.692	0.000

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT MR5: <i>MEDICAL RADIOLOGICAL DEFENSE (SDD)</i>
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Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
Remarks													
			Total Prior Years Cost	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract			
Project Cost Totals			0.802	8.276	1.143	0.000	1.143	0.000	10.221	0.000			

Remarks

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT MR5: <i>MEDICAL RADIOLOGICAL DEFENSE (SDD)</i>
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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015			
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
** MRADC - MRADC - Milestone B							■																					
MRADC - Non Clinical Animal Efficacy Studies								■	■	■	■	■	■	■	■	■												
MRADC - Process Development & cGMP Manufacturing Requirements						■	■	■	■	■	■	■	■	■	■	■												
MRADC - Product Formulation, Storage, and Delivery System on candidate 2						■	■	■	■	■	■	■	■	■														
MRADC - BLA Submission																■												
MRADC - FDA Approval																				■								
MRADC - Milestone C																								■				

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT MR5: <i>MEDICAL RADIOLOGICAL DEFENSE (SDD)</i>
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Schedule Details

Event	Start		End	
	Quarter	Year	Quarter	Year
** MRADC - MRADC - Milestone B	3	2010	3	2010
MRADC - Non Clinical Animal Efficacy Studies	4	2010	4	2012
MRADC - Process Development & cGMP Manufacturing Requirements	2	2010	4	2012
MRADC - Product Formulation, Storage, and Delivery System on candidate 2	2	2010	3	2012
MRADC - BLA Submission	2	2013	2	2013
MRADC - FDA Approval	2	2014	2	2014
MRADC - Milestone C	3	2014	3	2014

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT TE5: <i>TEST & EVALUATION (SDD)</i>
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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
TE5: <i>TEST & EVALUATION (SDD)</i>	37.444	36.593	15.901	0.000	15.901	12.243	4.238	14.614	15.300	Continuing	Continuing
Quantity of RDT&E Articles											

A. Mission Description and Budget Item Justification

This funding supports the Joint Project Manager Nuclear, Biological, Chemical Contamination Avoidance Product Director, Test Equipment, Strategy, and Support (PD TESS) efforts. PD TESS provides test infrastructure products for testing and evaluating chemical and biological defense systems throughout the life cycle acquisition process in support of the Milestone Decision Authority, Joint Project Managers, and the Test and Evaluation (T&E) community. PD TESS test infrastructure products are aligned in four groups to include: (1) Chemical Laboratory (Sense); (2) Biological Laboratory (Sense); (3) Field Simulant (Sense); (4) Individual Protection, Collective Protection and Decontamination (Shield and Sustain).

(1) Chemical Laboratory (Sense): Products for this area include a Non-Traditional Agent (NTA) Test Facility and a Dynamic Test Chamber (DTC) for chemical point sensors. The Dynamic Test Chamber provides a new capability for testing chemical point detection systems against chemical warfare agents in various environmental conditions. Major CBDP acquisition programs supported are: the Joint Chemical Agent Detector (JCAD).

(2) Sense Laboratory (Biological): Products for this area include a Whole System Live Agent Test (WSLAT) "Strung Out" Chamber; WSLAT "Full System" Chamber; and upgrade of a bio-level 3 facility located at Dugway Proving Ground (DPG). The WSLAT "Strung Out" Chamber supports Joint Biological Point Detection component testing in biological live agent environments. The WSLAT "Full System" Chamber supports testing of all biological detection systems in production configuration in biological live agent environments. The Baker Laboratory Upgrade will provide a bio-level 3 fabricated infrastructure to host the WSLAT "Full System" Chamber. The upgrade will include bio-level 3 support laboratories and analytical instrumentation. Major CBDP acquisition programs supported are: the Joint Biological Point Detection System (JBPDS)/JBPDS Block II; the Joint Biological Tactical Detection System (JBTDSD); and the Joint Biological Standoff Detection System (JBSDS) Block II.

(3) Field Simulant (Sense): Products for this area include a fully instrumented Simulant Test Grid and characterization of the Active Standoff Chamber (ASC) facilities. The Test Grid effort provides a fully instrumented 20 km by 40 km field simulant test capability that integrates cloud tracking equipment, meteorological equipment, test data network, C4ISR network, and operations center. The ASC effort provides a controlled simulant cloud characterization and facility. Major acquisition programs supported are: the Joint Chemical Agent Detector (JCAD); the Joint NBC Reconnaissance System (JNBCRS); the Joint Warning and Reporting Network (JWARN); the Joint Biological Standoff Detection System (JBSDS); the Joint Biological Point Detection System (JBPDS); the Joint Biological Tactical Detection System (JBTDSD); the Joint Effects Model (JEM); the Joint Operational Effects Federation (JOEF); and the Joint Expeditionary Collective Protection (JECPP) System.

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT TE5: <i>TEST & EVALUATION (SDD)</i>
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(4) Individual Protection, Collective Protection and Decontamination (Shield and Sustain): Products for this area include: a Small Item Decontamination Chamber; Individual Protection Ensemble (IPE) Mannequin; Man-in-Simulant Test (MIST) instrumentation; Individual Protection Equipment (IPE) Grid; Collective Protection (ColPro) instrumentation and facilities. The Small Item Decontamination Chamber provides an enhanced ability to conduct decontamination and residual agent off-gassing testing. The IPE Mannequin provides an articulated robotic mannequin that simulates Warfighters activities and includes under ensemble agent sensing capability for evaluating IPE against chemical warfare agents. The Man-in-Simulant Test instrumentation provides a near real time simulant sensor system to monitor penetration of simulant. The Individual Protection Equipment (IPE) Grid provides test procedures to establish commonality measurements for system level IPE performance tests. Chemical, Biological Agent Resistance Test (CBART) fixture provides a near real time testing capability under a range of environmental conditions for individual and collective protection materials. Collective Protection instrumentation and fixture upgrades provide improved test capabilities at Dugway Proving Ground, Dahlgren Naval Surface Warfare Center, and the Edgewood Chemical Biological Center for the evaluation of entire ColPro systems, subsystems and individual components. Acquisition Programs supported are: Joint Platform Interior Decontamination/Joint Material Decontamination System (JPID/JMDS); Joint Service Transportable Decontamination System (JSTDS); Joint Expeditionary Collective Protection (JECPC); Joint Collective Protection Equipment (JCPE); Protective Clothing; Joint Protective Aircrew Ensemble (JPACE); Joint Service General Purpose Mask (JSGPM); Joint Service Aircrew Mask (JSAM); and the Joint Chemical Ensemble (JCE).

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) PD TESS - IPE Mannequin <i>FY 2010 Plans:</i> Initiate IPE Mannequin and Chamber system design, fabrication and installation. <i>FY 2011 Base Plans:</i> Complete installation and conduct verification and validation testing.	0.000	13.200	6.300	0.000	6.300
2) PD TESS - IPE Man-in-Simulant Test (MIST) Upgrade <i>FY 2010 Plans:</i> Procure, verify and validate real-time MIST sensors.	0.000	0.659	0.000	0.000	0.000
3) PD TESS - Dynamic Test Chamber	4.610	0.750	0.000	0.000	0.000

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT TE5: <i>TEST & EVALUATION (SDD)</i>				
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2009 Accomplishments:</i> Completed assembly and installation and conducted validation testing.</p> <p><i>FY 2010 Plans:</i> Complete validation of the Dynamic Test Chamber.</p>						
<p>4) PD TESS - Test Grid Instrumentation Network & Design</p> <p><i>FY 2009 Accomplishments:</i> Procured chemical dissemination and referee instrumentation, installed test data network, and initiated build of the operations center.</p> <p><i>FY 2010 Plans:</i> Complete IOC for chemical simulant testing. Procure data network software build and operational demonstration. Procure operational demonstration hardware.</p> <p><i>FY 2011 Base Plans:</i> Conduct characterization testing of Biological referee instrumentation.</p>		23.175	14.441	4.530	0.000	4.530
<p>5) PD TESS - Decon Facility Upgrade</p> <p><i>FY 2009 Accomplishments:</i> Completed assembly and installation of small item decon chamber.</p>		0.193	0.000	0.000	0.000	0.000
<p>6) PD TESS - Whole System Live Agent Test (WSLAT)</p> <p><i>FY 2009 Accomplishments:</i> Continued construction of Baker Laboratory and procured referee instrumentation. Awarded WSLAT design, fabrication, and build contract.</p>		7.203	5.010	1.915	0.000	1.915

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2010 Plans:</i> Complete WSLAT design and initiate build. <i>FY 2011 Base Plans:</i> Complete WSLAT build and installation.						
7) PD TESS - ColPro Upgrades <i>FY 2009 Accomplishments:</i> Initiated biological and mechanical filtration for ColPro facilities.		0.528	0.000	0.000	0.000	0.000
8) PD TESS <i>FY 2009 Accomplishments:</i> Provided system engineering support for the design, build, integration and validation of test infrastructure products. <i>FY 2010 Plans:</i> Continue system engineering support for the design, build, integration and validation of test infrastructure products. <i>FY 2011 Base Plans:</i> Continue system engineering support for the design, build, integration and validation of test infrastructure products.		1.735	0.797	3.156	0.000	3.156
9) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.		0.000	0.461	0.000	0.000	0.000
Accomplishments/Planned Programs Subtotals		37.444	35.318	15.901	0.000	15.901

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010
Congressional Add: 1) Real Time Monitoring of Chemical Agents <i>FY 2010 Plans:</i> Congressional Interest Item - Real Time Monitoring of Chemical Agents, Chemical Agent Stimulants and Toxic Industrial Chemicals.	0.000	1.275
Congressional Adds Subtotals	0.000	1.275

C. Other Program Funding Summary (\$ in Millions)

<u>Line Item</u>	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011</u> <u>Base</u>	<u>FY 2011</u> <u>OCO</u>	<u>FY 2011</u> <u>Total</u>	<u>FY 2012</u>	<u>FY 2013</u>	<u>FY 2014</u>	<u>FY 2015</u>	<u>Cost To</u> <u>Complete</u>	<u>Total Cost</u>
• TE7: <i>TEST & EVALUATION (OP SYS DEV)</i>	7.037	4.870	4.813		4.813	4.779	4.750	5.660	5.615	Continuing	Continuing

D. Acquisition Strategy

PD TESS

The PD TESS program provides for the development and acquisition of new and enhanced test infrastructure to support the sense, shield, shape, and sustain mission areas for the Joint Service Chemical and Biological Defense Program (CBDP). The efforts are supported through competitive contract actions, academia, and other Government agencies. Infrastructure solutions will leverage commercially available systems to provide state-of-the-art capabilities that address current and future CBDP test and evaluation needs.

E. Performance Metrics

N/A

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT TE5: <i>TEST & EVALUATION (SDD)</i>
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Product Development (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** PD TESS - HW S - IPE Mannequin System Fabricate/Install	C/FFP	Midwest Research Institute Kansas City, MO	0.000	13.200	Jan 2010	6.300		0.000		6.300	0.000	19.500	0.000
HW S - WSLAT Chamber Fabrication/ Installation	C/FFP	TBD	4.521	5.010		1.915	Jan 2011	0.000		1.915	0.000	11.446	0.000
HW S - Test Grid Referee Instrumentation, Data Network and C4ISR	C/FFP	Lockheed Martin Integrated Systems Wall, NJ	40.959	14.441	Jan 2010	4.530		0.000		4.530	0.000	59.930	0.000
HW S - Dynamic Test Chamber Fabrication/ Installation	Reqn	NAVSEA (JHU-APL) Washington, DC	7.490	0.750		0.000		0.000		0.000	0.000	8.240	0.000
Congressional Interest Item - Real Time Monitoring of Chemical Agents, Chemical Agent Stimulants and Toxic Industrial Chemicals.	C/CPFF	TBD	0.000	1.275	Apr 2010	0.000		0.000		0.000	0.000	1.275	0.000
Subtotal			52.970	34.676		12.745		0.000		12.745	0.000	100.391	0.000

Remarks

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** PD TESS - OTH S - IPE MIST Validation	MIPR	Various	0.000	0.659	Jul 2010	0.000		0.000		0.000	0.000	0.659	0.000
Subtotal			0.000	0.659		0.000		0.000		0.000	0.000	0.659	0.000

Remarks
Test efforts are for the validation of capabilities.

Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** PD TESS - PM/MS S - Program Management/ Systems Engineering Support	MIPR	JPM NBCCA APG, MD	10.642	0.797	Oct 2009	3.156	Oct 2010	0.000		3.156	0.000	14.595	0.000
** ZSBIR - SBIR/STTR - Aggregated from ZSBIR-SBIR/STTR	MIPR	HQ AMC, Alexandria	0.000	0.461		0.000		0.000		0.000	0.000	0.461	0.000
Subtotal			10.642	1.258		3.156		0.000		3.156	0.000	15.056	0.000

Remarks

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT TE5: <i>TEST & EVALUATION (SDD)</i>

	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015							
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4				
** PD TESS - IPE MIST Sensors/Installations					■	■	■	■																								
Test Grid	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
WSLAT Chamber Design/Fabrication/Validation	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■												
Baker Laboratory Upgrade	■	■	■	■	■	■																										
Bio Standoff Facility Design/Fabrication/Installation					■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
DTC Fabrication/Installation	■	■	■	■	■	■	■																									
IPE Mannequin Design, Build, Install	■	■	■	■	■	■	■	■	■	■	■	■	■																			

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 5: <i>Development & Demonstration (SDD)</i>	R-1 ITEM NOMENCLATURE PE 0604384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (SDD)</i>	PROJECT TE5: <i>TEST & EVALUATION (SDD)</i>

Schedule Details

Event	Start		End	
	Quarter	Year	Quarter	Year
** PD TESS - IPE MIST Sensors/Installations	1	2010	4	2010
Test Grid	1	2009	4	2015
WSLAT Chamber Design/Fabrication/Validation	1	2006	4	2013
Baker Laboratory Upgrade	1	2008	2	2010
Bio Standoff Facility Design/Fabrication/Installation	2	2010	4	2015
DTC Fabrication/Installation	1	2008	3	2010
IPE Mannequin Design, Build, Install	1	2009	1	2012

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Exhibit R-2, RDT&E Budget Item Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 6: <i>RDT&E Management Support</i>	R-1 ITEM NOMENCLATURE PE 0605384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (RDT&E MGT SUPPORT)</i>
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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
Total Program Element	100.470	106.033	120.995	0.000	120.995	127.666	134.162	129.575	126.792	Continuing	Continuing
DT6: <i>JOINT DOCTRINE AND TRAINING SUPPORT (RDT&E MGT SUPPORT)</i>	5.388	6.637	6.332	0.000	6.332	5.547	5.722	5.889	6.004	Continuing	Continuing
DW6: <i>MAJOR RANGE AND TEST FACILITY BASE (MRTFB)</i>	53.710	54.461	60.274	0.000	60.274	67.308	73.054	74.610	74.332	Continuing	Continuing
LS6: <i>LABORATORY SUPPORT</i>	5.380	10.296	18.945	0.000	18.945	20.123	20.091	12.672	9.228	Continuing	Continuing
MS6: <i>RDT&E MGT SUPPORT</i>	30.144	27.424	29.714	0.000	29.714	29.711	30.233	31.193	31.917	Continuing	Continuing
O49: <i>JOINT CONCEPT DEVELOPMENT AND EXPERIMENTATION PROGRAM</i>	5.848	7.215	5.730	0.000	5.730	4.977	5.062	5.211	5.311	Continuing	Continuing

A. Mission Description and Budget Item Justification

This Budget Activity includes research, development, testing and evaluation management support for the Department of Defense (DoD) Chemical and Biological Defense Program (CBDP) and includes the CBDP Small Business Innovative Research (SBIR) program.

Program Element 0605384BP supports Joint Doctrine and Training (Project DT6), sustains the technical test capability at Dugway Proving Ground (DPG) (Project DW6); sustains the core Department of Defense (DoD) Science and Technology (S&T) laboratory infrastructure (Project LS6), provides for program management and financial management support (Project MS6), and supports the Joint Concept Development and Experimentation (JCDE) program (Project O49).

The Joint Training and Doctrine Support (DT6) project funds development of Joint Doctrine and Tactics, Techniques, and Procedures for developing CB defense systems. This project also funds CB modeling and simulation to support the Warfighter.

The Major Range and Test Facility Base (MRTFB) is a set of test installations, facilities, and ranges which are regarded as "national assets." These assets are sized, operated, and maintained primarily for DoD test and evaluation missions. However, the MRTFB facilities and ranges are also available to commercial and other users on a reimbursable basis. Dugway Proving Ground is designated as the primary element of the MRTFB to primarily conduct CB Defense test and evaluation. The DW6 Project provides operating funds to Dugway Proving Ground in accordance with the National Defense Authorization Act of 2003 (P/L/ 107-314 - sect 232) to ensure

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Exhibit R-2, RDT&E Budget Item Justification: PB 2011 Chemical and Biological Defense Program	DATE: February 2010
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APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 6: <i>RDT&E Management Support</i>	R-1 ITEM NOMENCLATURE PE 0605384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (RDT&E MGT SUPPORT)</i>
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that DOD test customers are only charged direct costs of testing and that overhead expenses are centrally funded. It finances the required institutional test operating costs. Institutional test operating costs include institutional civilian and contractor labor; repair and maintenance of test instrumentation, equipment, and facilities; and replacement of test equipment.

The Laboratory Support (LS6) project funds laboratory infrastructure to maintain and enhance DoD infrastructure capabilities to counter an expanding threat space, exploit advances in technology and develop and transition CB defense equipment and countermeasures to the Warfighter.

The management support (MS6) project, provides management support for the DoD CBDP to allow program overview and integration of overall medical and non-medical programs by the Assistant to the Secretary of Defense for Nuclear, Chemical, and Biological Defense Programs (ATSD(NCB)), through the Special Assistant, Chemical Biological Defense and Chemical Demilitarization Programs (SA(CBD&CDP)); execution management by the Defense Threat Reduction Agency (DTRA); integration of Joint requirements, management of training and doctrine by the Joint Requirements Office (JRO); Joint RDA planning, input to the Annual Report to Congress and Program Objective Memorandum (POM) development by the Program Analysis and Integration Office (PA&IO); review of Joint plans and the consolidated CB Defense POM Strategy by Army in its Executive Agent role.

The management support project also funds the Joint Test Infrastructure Working Group (JTIWG) program to execute the Test and Evaluation (T&E) Executive mission to establish test infrastructure investment strategy and adequate testing for Developmental Testing (DT) and Operational Testing (OT) of Department of Defense (DoD) Chemical Biological Defense (CBD) systems and components throughout the systems' acquisition life cycle, as required in the RDA Plan. The JTIWG program funds T&E Early Involvement, test threat planning, and T&E Standards planning and development to support testing the CBD systems for all services.

The Joint Concept Development and Experimentation (O49) project funds the planning, conduct, evaluation, and reporting on Joint tests (for other than developmental hardware) and accomplishment of operational research assessments in response to requirements received from the Services and the Combatant Commanders for already fielded equipment and systems.

This Budget Activity also funds Program Element 0605502BP, which supports the Small Business Innovative Research (SBIR) program. The overall objective of the Chemical and Biological Defense (CBD) SBIR program is to improve the transition or transfer of innovative CBD technologies between DoD components and the private sector for mutual benefit. The CBD program includes those technology efforts that maximize a strong defensive posture in a CB environment using passive and active means as deterrents. These technologies include CB detection; information assessment (identification, modeling, and intelligence); contamination avoidance; and protection of both individual soldiers and equipment.

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Exhibit R-2, RDT&E Budget Item Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY	R-1 ITEM NOMENCLATURE
0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i>	PE 0605384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (RDT&E MGT SUPPORT)</i>
BA 6: <i>RDT&E Management Support</i>	

B. Program Change Summary (\$ in Millions)

	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011 Base</u>	<u>FY 2011 OCO</u>	<u>FY 2011 Total</u>
Previous President's Budget	99.811	106.477	0.000	0.000	0.000
Current President's Budget	100.470	106.033	120.995	0.000	120.995
Total Adjustments	0.659	-0.444	120.995	0.000	120.995
• Congressional General Reductions		-0.444			
• Congressional Directed Reductions		0.000			
• Congressional Rescissions	0.000	0.000			
• Congressional Adds		0.000			
• Congressional Directed Transfers		0.000			
• Reprogrammings	1.812	0.000			
• SBIR/STTR Transfer	-1.153	0.000			
• Other Adjustments	0.000	0.000	120.995	0.000	120.995

Change Summary Explanation

Funding: N/A - Adjustments less than 10% of total program.

Schedule: N/A

Technical: N/A

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 6: <i>RDT&E Management Support</i>	R-1 ITEM NOMENCLATURE PE 0605384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (RDT&E MGT SUPPORT)</i>	PROJECT DT6: <i>JOINT DOCTRINE AND TRAINING SUPPORT (RDT&E MGT SUPPORT)</i>
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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
DT6: <i>JOINT DOCTRINE AND TRAINING SUPPORT (RDT&E MGT SUPPORT)</i>	5.388	6.637	6.332	0.000	6.332	5.547	5.722	5.889	6.004	Continuing	Continuing
Quantity of RDT&E Articles											

A. Mission Description and Budget Item Justification

The activities of this project directly support the Joint Service CB defense program; in particular, the development of Joint Chemical, Biological, Radiological, and Nuclear (CBRN) defense capability requirements and the improvement of CBRN defense related doctrine, education, training, and awareness at the Joint and Service levels. This effort provides for: (1) Development, coordination, and integration of Joint CBRN defense capability requirements; (2) Development/revision of medical and non-medical CBRN defense Multi-Service Tactics, Techniques, and Procedures (MTTP), Joint Doctrine and Tactics, Techniques, and Procedures (JTTP); (3) The CBRN Joint Senior Leader Course (JSLC); (4) Assistance in correcting training and doctrine deficiencies covered in the lessons learned process, combat operations, capability development studies and Department of Defense Inspector General (DODIG) and Government Accountability Office (GAO) reports; (5) Support of current and planned CBRN defense studies, analysis, training, exercises, and wargames; determine overlaps, duplication, and shortfalls; and build and execute programs to correct shortfalls in all aspects of CBRN defense across all DoD mission areas.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) JRO DT <i>FY 2009 Accomplishments:</i> Supported the revision and development of CBRN defense medical and physical sciences MTTPs. Supported the integration of CBRN defense considerations during the revision and development of selected Joint doctrine and JTTPs.	5.388	6.556	6.332	0.000	6.332

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 6: <i>RDT&E Management Support</i>	R-1 ITEM NOMENCLATURE PE 0605384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (RDT&E MGT SUPPORT)</i>	PROJECT DT6: <i>JOINT DOCTRINE AND TRAINING SUPPORT (RDT&E MGT SUPPORT)</i>				
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2010 Plans:</i> Continue to support the revision and development of CBRN defense medical and physical sciences MTTPs. Continue to support the integration of CBRN defense considerations during the revision and development of selected Joint doctrine and JTTPs.</p> <p><i>FY 2011 Base Plans:</i> Continue to support the revision and development of CBRN defense medical and physical sciences MTTPs. Continue to support the integration of CBRN defense considerations during the revision and development of selected Joint doctrine and JTTPs.</p>						
2) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.		0.000	0.081	0.000	0.000	0.000
Accomplishments/Planned Programs Subtotals		5.388	6.637	6.332	0.000	6.332
C. Other Program Funding Summary (\$ in Millions) N/A						
D. Acquisition Strategy N/A						
E. Performance Metrics N/A						

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY				R-1 ITEM NOMENCLATURE				PROJECT			
0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 6: <i>RDT&E Management Support</i>				PE 0605384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (RDT&E MGT SUPPORT)</i>				DW6: <i>MAJOR RANGE AND TEST FACILITY BASE (MRTFB)</i>			
COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
DW6: <i>MAJOR RANGE AND TEST FACILITY BASE (MRTFB)</i>	53.710	54.461	60.274	0.000	60.274	67.308	73.054	74.610	74.332	Continuing	Continuing
Quantity of RDT&E Articles											

A. Mission Description and Budget Item Justification

Project provides the technical capability for testing Department of Defense (DoD) Chemical and Biological (CB) defense materiel, equipment, and systems from concept through production at Dugway Proving Ground (DPG), a Major Range and Test Facility Base (MRTFB). Funding reflects compliance with National Defense Authorization Act (NDAA) for FY 2003 (Public Law 107-314 - December 2002), Sec 232, requiring Major Range and Test Facility Bases to be fully funded and that DoD test customers be charged for direct costs only.

DPG, a MRTFB, is the reliance center for all DoD CB defense testing and provides the United States' only combined range, chamber, toxic chemical lab, and bio-safety level three test facility. Total institutional test operating costs are to be provided by the Service component IAW DoD 3200.11.

DPG uses state-of-the-art chemical and life sciences test facilities and test chambers to perform CB defense testing of protective gear, decontamination systems, detectors, and equipment while totally containing chemical agents and biological pathogens. DPG also provides a fully instrumented outdoor range capability for testing with simulants that can be correlated to the laboratory testing with live agents.

Projects programmed for testing at DPG include: Joint Nuclear, Biological, and Chemical Reconnaissance System (JNBCRS); Joint Chemical Agent Detector (JCAD); Joint Biological Point Detection System (JBPDS); Joint Biological Stand-off Detection System (JBSDS); Joint Biological Agent Identification and Detection System (JBAIDS); Joint Biological Tactical Detection System (JBTDS); Whole System Live Agent Test (WSLAT); Joint Service Lightweight Integrated Suit Technology (JSLIST) Additional Sources Qualification 2 (JASQ 2); JSLIST Block II Glove Upgrade - Flame Resistant; JSLIST Performance Enhancement (JPE); JSLIST Combat Vehicle Crewman Coverage (JC3); Joint Protective Aircrew Ensemble (JPACE); Monitoring and Survey Sets, Kits, and Outfits (MSSKO); Joint Contaminated Surface Detector (JCSD); Joint Chemical, Biological and Radiological Agent Water Monitor (JCBRAWM); Nuclear, Biological and Chemical Environment Personal Hydration System (NEPHS); Analytical Lab System (ALS); Joint Expeditionary Collective Protection (JECP); Chemical Biological Protective Shelter (CBPS); Joint Service Aircrew Mask (JSAM); Joint Service Family of Decontamination Systems (JSFDS); Joint Materiel Decontamination System (JMDS); Human Remains Decontamination System (HRDS); Joint Warning and Reporting Network (JWARN); Joint Effects Model (JEM); and Joint Operational Effects Federation (JOEF).

B. Accomplishments/Planned Program (\$ in Millions)

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 6: <i>RDT&E Management Support</i>	R-1 ITEM NOMENCLATURE PE 0605384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (RDT&E MGT SUPPORT)</i>	PROJECT DW6: <i>MAJOR RANGE AND TEST FACILITY BASE (MRTFB)</i>				
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
5) DPG MRTFB <i>FY 2011 Base Plans:</i> Initiate Technology and Integration study and initial design work for Non Traditional Agent (NTA) activities.		0.000	0.000	0.500	0.000	0.500
6) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.		0.000	0.665	0.000	0.000	0.000
Accomplishments/Planned Programs Subtotals		53.710	54.461	60.274	0.000	60.274
C. Other Program Funding Summary (\$ in Millions) N/A						
D. Acquisition Strategy N/A						
E. Performance Metrics N/A						

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 6: <i>RDT&E Management Support</i>	R-1 ITEM NOMENCLATURE PE 0605384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (RDT&E MGT SUPPORT)</i>	PROJECT LS6: <i>LABORATORY SUPPORT</i>
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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
LS6: <i>LABORATORY SUPPORT</i>	5.380	10.296	18.945	0.000	18.945	20.123	20.091	12.672	9.228	Continuing	Continuing
Quantity of RDT&E Articles											

A. Mission Description and Budget Item Justification

This project (LS6) provides for the maintenance and enhancement of the DoD laboratory infrastructure capabilities to counter an expanding threat space, exploit advances in technology, and develop and transition chemical and biological (CB) defense equipment and countermeasures to the Warfighter. This laboratory infrastructure project upgrades key systems to the current state-of-the-art capabilities. Key systems include: gas filters, controls, emergency, mechanical/electrical, and structural systems. This project will ensure that the necessary surety operations can be conducted effectively and safely in support of Chemical and Biological Defense Program (CBDP) RDTE programs. As a force multiplier, this project will provide more robust capabilities to the CBDP and ensure continuity of operations and environmental compliance.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) LABINF - Gas Filters <i>FY 2009 Accomplishments:</i> Modernized existing gas filters to include developing new filter designs with the capability of protecting against emerging threat agents. <i>FY 2010 Plans:</i> Modernize existing gas filters to include developing new filter designs with the capability of protecting against emerging threat agents. <i>FY 2011 Base Plans:</i> Modernize existing gas filters to include developing new filter designs with the capability of protecting against emerging threat agents.	1.200	1.245	1.314	0.000	1.314

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 6: <i>RDT&E Management Support</i>		R-1 ITEM NOMENCLATURE PE 0605384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (RDT&E MGT SUPPORT)</i>		PROJECT LS6: <i>LABORATORY SUPPORT</i>		
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
2) LABINF - Control Systems <i>FY 2009 Accomplishments:</i> Modernized mechanical and pneumatic control systems to full digital controls. <i>FY 2010 Plans:</i> Modernize mechanical and pneumatic control systems to full digital controls. <i>FY 2011 Base Plans:</i> Modernize mechanical and pneumatic control systems to full digital controls.		0.991	0.995	0.896	0.000	0.896
3) LABINF - Emergency Systems <i>FY 2009 Accomplishments:</i> Modernized emergency systems to increase reliability and safety. <i>FY 2010 Plans:</i> Modernize emergency systems to increase reliability and safety. <i>FY 2011 Base Plans:</i> Modernize emergency systems to increase reliability and safety.		0.992	0.900	0.920	0.000	0.920
4) LABINF - Mechanical/Electrical Systems <i>FY 2009 Accomplishments:</i> Provided upgrades to key systems to ensure worker safety, environmental compliance, and continuity of operations. Upgrades included low-flow hood alarms, redundant exhaust fans and HVAC controllers.		1.205	1.279	1.254	0.000	1.254

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010				
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 6: <i>RDT&E Management Support</i>		R-1 ITEM NOMENCLATURE PE 0605384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (RDT&E MGT SUPPORT)</i>		PROJECT LS6: <i>LABORATORY SUPPORT</i>				
B. Accomplishments/Planned Program (\$ in Millions)								
				FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2010 Plans:</i> Provides key chemical and biological defense effort upgrades, initial outfitting, and equipment for the United States Army Medical Research Institute of Infectious Diseases (USAMRIID) and United States Army Medical Research Institute of Chemical Defense (USAMRICD) infrastructure.</p> <p><i>FY 2011 Base Plans:</i> Provides key chemical and biological defense effort upgrades, initial outfitting, and equipment for the USAMRIID and USAMRICD infrastructure.</p>								
7) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.				0.000	0.125	0.000	0.000	0.000
Accomplishments/Planned Programs Subtotals				5.380	10.296	18.945	0.000	18.945
C. Other Program Funding Summary (\$ in Millions)								
N/A								
D. Acquisition Strategy								
N/A								
E. Performance Metrics								
N/A								

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 6: <i>RDT&E Management Support</i>	R-1 ITEM NOMENCLATURE PE 0605384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (RDT&E MGT SUPPORT)</i>	PROJECT MS6: <i>RDT&E MGT SUPPORT</i>
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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
MS6: <i>RDT&E MGT SUPPORT</i>	30.144	27.424	29.714	0.000	29.714	29.711	30.233	31.193	31.917	Continuing	Continuing
Quantity of RDT&E Articles											

A. Mission Description and Budget Item Justification

This project provides management support for the DoD CBDP. It includes program oversight and integration of overall medical and non-medical programs by the Assistant to the Secretary of Defense for Nuclear and Chemical and Biological Defense Programs (ATSD(NCB)) defense programs through the Deputy Assistant to the Secretary of Defense for Chemical and Biological Defense/Chemical Demilitarization (ODATSD(CBD/CD)). Funds execution management is provided by DTRA.

The project also provides for the development, coordination and integration of Joint Chemical, Biological, Radiological and Nuclear (CBRN) defense capability requirements, including assistance and support to the Combatant Commanders and Services to improve CBRN defense related doctrine, education, training, and awareness by the Joint Requirements Office (JRO) Joint CBRN Defense Research, Development, and Acquisition (RDA) planning; and input to the CBD Annual Report to Congress, and program guidance development by the Program Analysis and Integration Office (PA&IO).

The project includes programming support for the Joint Service CB Information System (JSCBIS) which serves as a budgetary and informational database for the DoD CBDP. Also included within the project is financial management services include fund distribution, execution reporting and fiscal financial statements.

This project also supports the Test and Evaluation (T&E) Executive, who is responsible for the planning, balancing, and oversight of test infrastructure and test technology requirements to support Developmental Testing (DT) and Operational Testing (OT) of DoD CBD systems, as outlined in the RDA Plan. The T&E Executive guides JPEO planning and coordination with the Operational Test Activities to plan a series of methodology, instrumentation, and associated validation efforts that provide test infrastructure and technologies for testing RDA systems needed to support all Services, and to ensure the adequacy of testing for RDA systems in alignment with acquisition schedules and associated decision points.

The CBDP T&E Executive directly supports OSD T&E oversight acquisition programs and provides the mechanism for early T&E involvement in the acquisition process. The CBDP T&E Executive provides the T&E infrastructure investment strategy and coordinates investment planning and T&E capabilities validation among the Joint Service Community to ensure that program needs are met.

B. Accomplishments/Planned Program (\$ in Millions)

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 6: <i>RDT&E Management Support</i>	R-1 ITEM NOMENCLATURE PE 0605384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (RDT&E MGT SUPPORT)</i>	PROJECT MS6: <i>RDT&E MGT SUPPORT</i>
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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2010 Plans:</i> Joint Test Infrastructure Working Group (JTIWG) - Continue Test and Evaluation (T&E) Executive mission support to ensure credible testing of Chemical Biological Defense Program (CBDP) systems and support to the Director for Operational Test and Evaluation (DOT&E) for OSD T&E Oversight. Continue direct support to the Joint Program Executive Office for Chemical Biological Defense (JPEO-CBD) and the Joint Requirements Office (JRO) Integrated Process Teams (IPTs) and Integrated Concept teams (ICTs) providing technical assistance to structure acquisition programs and test scopes. Continue early involvement of the Operational Test Agencies (OTAs) and other T&E organizations in T&E infrastructure planning. Continue development of threat test support documentation to support developmental and operational tests in which an operational threat must be realistically presented, including Joint Chemical Agent Detector (JCAD), Joint Biological Standoff Detection System (JBSDS), Joint Biological Tactical Detection System (JBTDSD), Joint Service Lightweight Nuclear, Biological, Chemical Reconnaissance System (JSLNBCRS), Joint Biological Agent Identification and Diagnostic System (JBAIDS), Joint Warning and Reporting Network (JWARN), Joint Service Transportable Decontamination System Small Scale (JSTDS-SS), and the Improved Point Detection System (IPDS). Continue support to JPEO-CBD and Joint Science and Technology Office (JSTO)-CB regarding specific test methodology and test technology needs, to include updates to the Technology Transition documents, participation in scientific review panels, and review of technology/methodology and development plans. Continue to provide guidance to improve the Test and Evaluation Master Plan (TEMP)s for acquisition programs, threat support documentation development, and development of T&E Capabilities Needs Statements and to expedite Lead OTA assignment and overall coordination. Continue to lead the International T&E methodology development and standardization efforts to support the Australia, Canadian, UK, and US Memorandum of Understanding (MOU). Provide T&E infrastructure input to the Program Objective Memorandum (POM) process and support JRO, Program Analysis and Integration Office (PA&IO), and SA(CBD & CDP) in development and defense of POM and Budget submissions.</p>					

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 6: <i>RDT&E Management Support</i>	R-1 ITEM NOMENCLATURE PE 0605384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (RDT&E MGT SUPPORT)</i>	PROJECT MS6: <i>RDT&E MGT SUPPORT</i>
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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2011 Base Plans:</i> Develop assessments to support RDA Planning. Provide analytic programmatic support for development of program guidance, the Program, Budget and Execution Reviews, and the PB submissions. Respond to specialized evaluation studies throughout the PPBE process. Provide JSCBIS database management.</p>					
<p>5) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.</p>	0.000	0.335	0.000	0.000	0.000
Accomplishments/Planned Programs Subtotals	30.144	27.424	29.714	0.000	29.714

C. Other Program Funding Summary (\$ in Millions)

N/A

D. Acquisition Strategy

N/A

E. Performance Metrics

N/A

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 6: <i>RDT&E Management Support</i>	R-1 ITEM NOMENCLATURE PE 0605384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (RDT&E MGT SUPPORT)</i>	PROJECT O49: <i>JOINT CONCEPT DEVELOPMENT AND EXPERIMENTATION PROGRAM</i>
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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
O49: <i>JOINT CONCEPT DEVELOPMENT AND EXPERIMENTATION PROGRAM</i>	5.848	7.215	5.730	0.000	5.730	4.977	5.062	5.211	5.311	Continuing	Continuing
Quantity of RDT&E Articles											

A. Mission Description and Budget Item Justification

The objectives of the Joint Concept Development and Experimentation (JCDE) program are to plan, conduct, evaluate, and report on joint tests and experiments (for other than developmental hardware) and accomplish capability development assessments. This program will provide ongoing input to the Combatant Commanders and Services for development of doctrine, policy, training procedures, and feedback into the Joint Capabilities Integration and Development System (JCID) and acquisition processes.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) JCDE <i>FY 2009 Accomplishments:</i> Supported the JCD for CBRND in conducting workshops, studies, war games and limited objective experiments to explore, refine, and validate potential solutions and alternatives that will update and improve the Joint CBRND concept. <i>FY 2010 Plans:</i> Support the JCD for CBRND in conducting workshops, studies, war games and limited objective experiments to explore, refine, and validate potential solutions and alternatives that will update and improve the Joint CBRND concept.	5.848	7.125	5.730	0.000	5.730

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010				
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 6: <i>RDT&E Management Support</i>		R-1 ITEM NOMENCLATURE PE 0605384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (RDT&E MGT SUPPORT)</i>		PROJECT O49: <i>JOINT CONCEPT DEVELOPMENT AND EXPERIMENTATION PROGRAM</i>				
B. Accomplishments/Planned Program (\$ in Millions)								
				FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2011 Base Plans:</i> Support the JCD for CBRND in conducting workshops, studies, war games and limited objective experiments to explore, refine, and validate potential solutions and alternatives that will update and improve the Joint CBRND concept.								
2) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.				0.000	0.090	0.000	0.000	0.000
Accomplishments/Planned Programs Subtotals				5.848	7.215	5.730	0.000	5.730
C. Other Program Funding Summary (\$ in Millions)								
N/A								
D. Acquisition Strategy								
N/A								
E. Performance Metrics								
N/A								

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Exhibit R-2, RDT&E Budget Item Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY			R-1 ITEM NOMENCLATURE								
0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 6: <i>RDT&E Management Support</i>			PE 0605502BP: <i>SMALL BUSINESS INNOVATIVE RESEARCH (SBIR)</i>								
COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
Total Program Element	12.713	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	Continuing	Continuing
SB6: <i>SMALL BUSINESS INNOVATIVE RESEARCH (SBIR)</i>	12.713	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	Continuing	Continuing

A. Mission Description and Budget Item Justification

The overall objective of the CBD SBIR program is to improve the transition or transfer of innovative CBD technologies between DoD components and the private sector for mutual benefit. The CBD program includes those technology efforts that maximize a strong defensive posture in a biological or chemical environment using passive and active means as deterrents. These technologies include chemical and biological detection; information assessment, which includes identification, modeling, and intelligence; contamination avoidance; and protection of both individual soldiers and equipment.

B. Program Change Summary (\$ in Millions)

	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011 Base</u>	<u>FY 2011 OCO</u>	<u>FY 2011 Total</u>
Previous President's Budget	0.000	0.000	0.000	0.000	0.000
Current President's Budget	12.713	0.000	0.000	0.000	0.000
Total Adjustments	12.713	0.000	0.000	0.000	0.000
• Congressional General Reductions		0.000			
• Congressional Directed Reductions		0.000			
• Congressional Rescissions	0.000	0.000			
• Congressional Adds		0.000			
• Congressional Directed Transfers		0.000			
• Reprogrammings	0.000	0.000			
• SBIR/STTR Transfer	12.713	0.000			
• Other Adjustments	0.000	0.000	0.000	0.000	0.000

Change Summary Explanation

Funding: FY09 - Funding transferred and applied to SBIR program (+\$12,713K).

Schedule: N/A

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Exhibit R-2, RDT&E Budget Item Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 6: <i>RDT&E Management Support</i>	R-1 ITEM NOMENCLATURE PE 0605502BP: <i>SMALL BUSINESS INNOVATIVE RESEARCH (SBIR)</i>
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Technical: N/A

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 6: <i>RDT&E Management Support</i>				R-1 ITEM NOMENCLATURE PE 0605502BP: <i>SMALL BUSINESS INNOVATIVE RESEARCH (SBIR)</i>				PROJECT SB6: <i>SMALL BUSINESS INNOVATIVE RESEARCH (SBIR)</i>			
COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
SB6: <i>SMALL BUSINESS INNOVATIVE RESEARCH (SBIR)</i>	12.713	0.000	0.000	0.000	0.000	0.000	0.000	0.000	0.000	Continuing	Continuing
Quantity of RDT&E Articles											

A. Mission Description and Budget Item Justification

The SBIR Program is a Congressionally mandated program established to increase the participation of small business in federal research and development (R&D). Currently, each participating government agency must reserve 2.5% of its extramural R&D for SBIR awards to competing small businesses. The goal of the SBIR Program is to invest in the innovative capabilities of the small business community to help meet government R&D objectives while allowing small companies to develop technologies and products which they can then commercialize through sales back to the government or in the private sector.

The Small Business Technology Transfer (STTR) Program like SBIR, is a Government-wide program, mandated by the Small Business Research and Development Enhancement Act of 1992, PL 102-564. STTR was established in FY94 as a three-year pilot program. In early 1996, the General Accounting Office (GAO) conducted a comprehensive review of the Government-wide STTR Program to determine the effectiveness of the pilot program. Upon review of the GAO report, Congress voted to reauthorize the STTR Program to the year 2000, consistent with the authorization period for the SBIR Program.

STTR was established as a companion program to the SBIR Program and is executed in essentially the same manner; however, there are several distinct differences. The STTR Program provides a mechanism for participation by university, Federally-Funded Research and Development Centers (FFRDCs), and other non-profit research institutions. Specifically, the STTR Program is designed to provide an incentive for small companies and research at academic institutions and non-profit research and development institutions to work together to move emerging technical ideas from the laboratory to the marketplace to foster high-tech economic development and to advance U.S. economic competitiveness. Each STTR proposal must be submitted by a team which includes a small business (as the prime contractor for contracting purposes) and at least one research institution, which have entered into a Cooperative Research and Development Agreement for the purposes of the STTR effort. Furthermore, the project must be divided up such that the small business performs at least 40% of the work and the research institution(s) performs at least 30% of the work. The remainder of the work may be performed by either party or a third party. The budget is separate from the SBIR budget and is significantly smaller (0.15% of the extramural R&D budget vs. 2.5% for the SBIR Program).

The DoD has consolidated management and oversight of the CBDP into a single office within the OSD. The Army was designated as the Executive Agent for coordination and integration of the Chemical and Biological Defense (CBD) program. The executive agent for the SBIR/STTR portion of the program is the Army Research Office-Washington.

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 6: <i>RDT&E Management Support</i>	R-1 ITEM NOMENCLATURE PE 0605502BP: <i>SMALL BUSINESS INNOVATIVE RESEARCH (SBIR)</i>	PROJECT SB6: <i>SMALL BUSINESS INNOVATIVE RESEARCH (SBIR)</i>
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The overall objective of the CBD SBIR/STTR program is to improve the transition or transfer of innovative CBD technologies between DoD components and the private sector for mutual benefit. The CBD program includes those technology efforts that maximize a strong defensive posture in a biological or chemical environment using passive and active means as deterrents. These technologies include chemical and biological detection; information assessment, which includes identification, modeling, and intelligence; contamination avoidance; and protection of both individual soldiers and equipment.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) ZSBIR <i>FY 2009 Accomplishments:</i> Small Business Innovative Research.	12.713	0.000	0.000	0.000	0.000
Accomplishments/Planned Programs Subtotals	12.713	0.000	0.000	0.000	0.000

C. Other Program Funding Summary (\$ in Millions)

N/A

D. Acquisition Strategy

N/A

E. Performance Metrics

N/A

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Exhibit R-2, RDT&E Budget Item Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 7: <i>Operational Systems Development</i>	R-1 ITEM NOMENCLATURE PE 0607384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (OP SYS DEV)</i>
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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
Total Program Element	12.494	6.172	6.634	0.000	6.634	9.317	10.639	14.763	11.873	Continuing	Continuing
IP7: <i>INDIVIDUAL PROTECTION OPERATIONAL SYS DEV</i>	4.560	0.000	0.000	0.000	0.000	2.869	4.371	7.548	4.681	Continuing	Continuing
IS7: <i>INFORMATION SYSTEMS (OP SYS DEV)</i>	0.897	1.302	1.821	0.000	1.821	1.669	1.518	1.555	1.577	Continuing	Continuing
TE7: <i>TEST & EVALUATION (OP SYS DEV)</i>	7.037	4.870	4.813	0.000	4.813	4.779	4.750	5.660	5.615	Continuing	Continuing

A. Mission Description and Budget Item Justification

This program element provides development efforts to upgrade systems in the Department of Defense (DoD) Chemical Biological Defense Program that have been fielded or have received approval for full rate production and anticipate production funding in the current or subsequent fiscal year.

Efforts in this program element support the upgrade of fielded CB defense equipment against emerging chemical threat agents and toxic industrial chemicals. Specifically this program supports (1) the upgrade of individual protection systems, (2) the JPEO-CBD Software Support Activity (SSA), and (3) revitalization and technical upgrade of existing instrumentation and equipment at Dugway Proving Ground (DPG).

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Exhibit R-2, RDT&E Budget Item Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY	R-1 ITEM NOMENCLATURE
0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i>	PE 0607384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (OP SYS DEV)</i>
BA 7: <i>Operational Systems Development</i>	

B. Program Change Summary (\$ in Millions)

	<u>FY 2009</u>	<u>FY 2010</u>	<u>FY 2011 Base</u>	<u>FY 2011 OCO</u>	<u>FY 2011 Total</u>
Previous President's Budget	12.640	6.198	0.000	0.000	0.000
Current President's Budget	12.494	6.172	6.634	0.000	6.634
Total Adjustments	-0.146	-0.026	6.634	0.000	6.634
• Congressional General Reductions		-0.026			
• Congressional Directed Reductions		0.000			
• Congressional Rescissions	0.000	0.000			
• Congressional Adds		0.000			
• Congressional Directed Transfers		0.000			
• Reprogrammings	0.000	0.000			
• SBIR/STTR Transfer	-0.146	0.000			
• Other Adjustments	0.000	0.000	6.634	0.000	6.634

Congressional Add Details (\$ in Millions, and Includes General Reductions)

Project: IP7: *INDIVIDUAL PROTECTION OPERATIONAL SYS DEV*

Congressional Add: 1) *JCEIII*

	<u>FY 2009</u>	<u>FY 2010</u>
	2.371	0.000
Congressional Add Subtotals for Project: IP7	2.371	0.000
Congressional Add Totals for all Projects	2.371	0.000

Change Summary Explanation

Funding: N/A - Adjustments less than 10% of total program.

Schedule: N/A

Technical: N/A

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program								DATE: February 2010			
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 7: <i>Operational Systems Development</i>				R-1 ITEM NOMENCLATURE PE 0607384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (OP SYS DEV)</i>				PROJECT IP7: <i>INDIVIDUAL PROTECTION OPERATIONAL SYS DEV</i>			
COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
IP7: <i>INDIVIDUAL PROTECTION OPERATIONAL SYS DEV</i>	4.560	0.000	0.000	0.000	0.000	2.869	4.371	7.548	4.681	Continuing	Continuing
Quantity of RDT&E Articles											
A. Mission Description and Budget Item Justification This project provides developmental efforts to upgrade fielded Individual Protection (IP) systems to include battle dress uniform, gloves, footwear and masks for protection against Non-Traditional Agents.											
B. Accomplishments/Planned Program (\$ in Millions)											
						FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	
1) JSGPM Filtration - <i>FY 2009 Accomplishments:</i> Initiated IPT to explore integration concepts.						1.639	0.000	0.000	0.000	0.000	
2) ESLI - <i>FY 2009 Accomplishments:</i> Conducted critical design review for End-of-Service Life Indicator (ESLI), and fabricated final prototype.						0.550	0.000	0.000	0.000	0.000	
Accomplishments/Planned Programs Subtotals						2.189	0.000	0.000	0.000	0.000	
						FY 2009	FY 2010				
Congressional Add: 1) JCEIII						2.371	0.000				

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 7: <i>Operational Systems Development</i>	R-1 ITEM NOMENCLATURE PE 0607384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (OP SYS DEV)</i>	PROJECT IP7: <i>INDIVIDUAL PROTECTION OPERATIONAL SYS DEV</i>
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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010
<p><i>FY 2009 Accomplishments:</i> Congressional Interest Item - JSGPM Filtration. Initiated IPT to explore integration concepts, develop new performance specification with increase Toxic Industrial Chemicals (TIC) requirements, start filter qualification. Awarded contract for new filtration development.</p>		
Congressional Adds Subtotals	2.371	0.000

C. Other Program Funding Summary (\$ in Millions)

N/A

D. Acquisition Strategy

LCBE

The Lightweight Chemical Biological Ensemble (LCBE) program will pursue an evolutionary incremental approach to provide capability to the Warfighter. Each increment of LCBE will provide technologies with military utility that are modular in function, and offer improvement in form and fit over current systems. The LCBE program will develop, integrate, test, procure and field systems that increase Warfighter operational performance in a CBRN environment via the use of emerging technologies and by leveraging tradespace in areas such as protection level, heat stress, durability, antimicrobial properties, launderability, self-detoxification, protection time, etc. Where appropriate, modeling and simulation tools will be used to lower LCBE program risks, reduce costs and ensure a high confidence in selected technologies.

LCBE INCREMENT 1

The LCBE will use an evolutionary acquisition strategy with phased development. The first LCBE increment will provide an operationally useful and supportable capability in as short a time as possible. Accordingly, Increment 1 of LCBE will incorporate an accelerated development cycle leveraging existing COTS technologies that will, at a minimum, provide a lightweight CB protective garment capability. Gate testing and down-selection of prototypes will comprise the initial phases of the Government's testing program. A competitively awarded contract is planned for DT and Operational Assessment (OA) will occur prior to MS C. Appropriate system requirements reviews, test readiness reviews, producibility reviews and audits will be scheduled as required prior to each milestone.

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 7: <i>Operational Systems Development</i>	R-1 ITEM NOMENCLATURE PE 0607384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (OP SYS DEV)</i>	PROJECT IP7: <i>INDIVIDUAL PROTECTION OPERATIONAL SYS DEV</i>
<p>Future increments of LCBE shall be defined via separate Capability Development Document (CDDs)/Capability Production Document (CPDs) and will follow a similar path/process from MS A or MS B through MS C/FRP and will leverage preceding efforts to the greatest extent possible, maintaining commonality and synergy across all increments.</p> <p><u>E. Performance Metrics</u> N/A</p>		

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 7: <i>Operational Systems Development</i>	R-1 ITEM NOMENCLATURE PE 0607384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (OP SYS DEV)</i>	PROJECT IP7: <i>INDIVIDUAL PROTECTION OPERATIONAL SYS DEV</i>
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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015							
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4				
** LCBE - Initiate IPT	■	■	■	■																												
Fabricate ESLI Prototype		■	■																													
ESLI Test & Evaluation		■	■	■	■																											

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 7: <i>Operational Systems Development</i>	R-1 ITEM NOMENCLATURE PE 0607384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (OP SYS DEV)</i>	PROJECT IP7: <i>INDIVIDUAL PROTECTION OPERATIONAL SYS DEV</i>
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Schedule Details

Event	Start		End	
	Quarter	Year	Quarter	Year
** LCBE - Initiate IPT	1	2009	4	2009
Fabricate ESLI Prototype	2	2009	3	2009
ESLI Test & Evaluation	2	2009	1	2010

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 7: <i>Operational Systems Development</i>				R-1 ITEM NOMENCLATURE PE 0607384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (OP SYS DEV)</i>				PROJECT IS7: <i>INFORMATION SYSTEMS (OP SYS DEV)</i>			
COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
IS7: <i>INFORMATION SYSTEMS (OP SYS DEV)</i>	0.897	1.302	1.821	0.000	1.821	1.669	1.518	1.555	1.577	Continuing	Continuing
Quantity of RDT&E Articles											

A. Mission Description and Budget Item Justification

The project supports the JPEO-CBD Software Support Activity (SSA). The JPEO-CBD SSA is a JPEO-CBD enterprise-wide, user developmental support and Service organization focusing on development assistance and net-centric interoperability. The SSA provides the CBRN Warfighter with Joint Service solutions for Information Assurance, Verification, Validation and Accreditation (VV&A), and Data Management; interoperable and integrated net-centric, Service-oriented, composable solutions for CBD; and infusion of latest technologies into programs of record. CBRN user community and related communities of interest have need for CBRN "plug and play" capability to allow interoperability and re-configurability across the enterprise. The requirement for net-centric, composable solutions provides the near term foundation for the Warfighter's ability to communicate his CBRN solutions and interoperate with other Service operational systems. It also supports a longer term ability to interoperate with related agencies and to reduce the Warfighter's CBRN footprint as technologies improve.

B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
1) SSA Integrated Architectures Support <i>FY 2009 Accomplishments:</i> Provided and updated program of record integrated architectures. <i>FY 2010 Plans:</i> Continue to provide and update program of record integrated architectures. <i>FY 2011 Base Plans:</i> Continue to provide and update program of record integrated architectures.	0.117	0.162	0.224	0.000	0.224
2) SSA Chemical, Biological, Radiological, Nuclear (CBRN) Data Model	0.108	0.153	0.244	0.000	0.244

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 7: <i>Operational Systems Development</i>	R-1 ITEM NOMENCLATURE PE 0607384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (OP SYS DEV)</i>	PROJECT IS7: <i>INFORMATION SYSTEMS (OP SYS DEV)</i>				
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p><i>FY 2009 Accomplishments:</i> Analyzed requirements and assisted programs with implementation of the CBRN data model.</p> <p><i>FY 2010 Plans:</i> Continue to analyze requirements and assist programs with implementation of the CBRN data model.</p> <p><i>FY 2011 Base Plans:</i> Continue to analyze requirements and assist programs with implementation of the CBRN data model.</p>						
<p>3) SSA CBRN Data Model Support</p> <p><i>FY 2009 Accomplishments:</i> Provided CBRN Data Model Reference implementations.</p> <p><i>FY 2010 Plans:</i> Continue to provide CBRN Data Model Reference implementations.</p> <p><i>FY 2011 Base Plans:</i> Continue to provide CBRN Data Model Reference implementations.</p>		0.083	0.162	0.228	0.000	0.228
<p>4) SSA CBRN Data Model Updates</p> <p><i>FY 2009 Accomplishments:</i> Supported CBRN Data Model updates.</p> <p><i>FY 2010 Plans:</i> Continue to support CBRN Data Model updates.</p> <p><i>FY 2011 Base Plans:</i> Continue to support CBRN Data Model updates.</p>		0.051	0.100	0.140	0.000	0.140

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 7: <i>Operational Systems Development</i>	R-1 ITEM NOMENCLATURE PE 0607384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (OP SYS DEV)</i>	PROJECT IS7: <i>INFORMATION SYSTEMS (OP SYS DEV)</i>				
B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
5) SSA Information Assurance Compliance Testing <i>FY 2009 Accomplishments:</i> Provided Information Assurance compliance testing for JPEO-CBD programs. <i>FY 2010 Plans:</i> Continue to provide Information Assurance compliance testing for JPEO-CBD programs. <i>FY 2011 Base Plans:</i> Continue to provide Information Assurance compliance testing for JPEO-CBD programs.		0.058	0.123	0.173	0.000	0.173
6) SSA Modeling and Simulation Support <i>FY 2009 Accomplishments:</i> Provided Modeling and Simulation IPT and Accreditation Steering Group support. <i>FY 2010 Plans:</i> Provide Modeling and Simulation IPT and Accreditation Steering Group support. <i>FY 2011 Base Plans:</i> Provide Modeling and Simulation IPT and Accreditation Steering Group support.		0.121	0.153	0.208	0.000	0.208
7) SSA Enterprise Information Support Plans <i>FY 2009 Accomplishments:</i> Provided Information Support Plan (ISP) development support for JPEO-CBD programs. <i>FY 2010 Plans:</i> Continue to provide ISP development support for JPEO-CBD programs.		0.116	0.179	0.248	0.000	0.248

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program				DATE: February 2010		
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B. Accomplishments/Planned Program (\$ in Millions)						
		FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<i>FY 2011 Base Plans:</i> Continue to provide ISP development support for JPEO-CBD programs.						
8) SSA Help Desk <i>FY 2009 Accomplishments:</i> Provided developmental Help Desk support for JPEO-CBD programs and users until they transitioned to sustainment funding.		0.075	0.000	0.000	0.000	0.000
9) SSA <i>FY 2009 Accomplishments:</i> Provided Net-Centric Policy implementation assistance. <i>FY 2010 Plans:</i> Continue to provide Net-Centric Policy implementation assistance. <i>FY 2011 Base Plans:</i> Continue to provide Net-Centric Policy implementation assistance.		0.066	0.087	0.123	0.000	0.123
10) SSA <i>FY 2009 Accomplishments:</i> Provided Common CBRN Sensor Interface (CCSI) reference implementation guidance. <i>FY 2010 Plans:</i> Provide CCSI reference implementation guidance. <i>FY 2011 Base Plans:</i> Provide CCSI reference implementation guidance.		0.064	0.084	0.118	0.000	0.118

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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
11) SSA <i>FY 2009 Accomplishments:</i> Supported CCSI updates. <i>FY 2010 Plans:</i> Support CCSI updates. <i>FY 2011 Base Plans:</i> Support CCSI updates.	0.038	0.083	0.115	0.000	0.115
12) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.	0.000	0.016	0.000	0.000	0.000
Accomplishments/Planned Programs Subtotals	0.897	1.302	1.821	0.000	1.821

C. Other Program Funding Summary (\$ in Millions)

N/A

D. Acquisition Strategy

SSA

The JPEO-CBD Software Support Activity (SSA) is a JPEO-CBD user support organization spanning and supporting all Joint Project Managers (JPMs) and JPEO-CBD Directorates. The SSA provides enterprise-wide services and coordination across all JPEO-CBD Programs of Record (PORs) that contain data or software, or are capable of linking to the Global Information Grid (GIG). The SSA facilitates interoperability, integration, and supportability of existing and developing IT and National Security Systems (NSS) across the JPEO and all JPMs.

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<p>Phase 1a identifies JPEO-CBD JPMs and programs that deal with data or software, and have an IT component. This will be followed by coordination with the JPMs and programs to facilitate the concepts of interoperability, integration and supportability of enterprise-wide services. Next follows work with user communities to develop and demonstrate enterprise-wide common architectures, products and services. [BA5 - System Development and Demonstration].</p> <p>Phase 1b established management and control measures for tracking and reporting progress of the various elements described in Phases 1 and 2. This includes establishing, tracking, and performing configuration management of inventories and databases of IT systems and their states of interoperability and information assurance compliance. [BA6 - RDT&E Management Support].</p> <p>Phase 2 will support the application of the enterprise-wide architectures, products and services into the programs, with verification of compliance with the defined products and services. [BA7 - Operational Systems Development].</p> <p><u>E. Performance Metrics</u> N/A</p>		

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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Product Development (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** SSA - Development Services	MIPR	SPAWAR System Center San Diego, CA	0.602	0.393	Oct 2009	0.609	Oct 2010	0.000		0.609	0.000	1.604	0.000
Subtotal			0.602	0.393		0.609		0.000		0.609	0.000	1.604	0.000

Remarks

Support (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** SSA - Develop Support Activities	MIPR	SPAWAR Systems Center San Diego, CA	0.555	0.408	Oct 2009	0.628	Oct 2010	0.000		0.628	0.000	1.591	0.000
Subtotal			0.555	0.408		0.628		0.000		0.628	0.000	1.591	0.000

Remarks

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** SSA - Integration Verification and Valuation (IV&V)	MIPR	SPAWAR Systems Center San Diego, CA	0.425	0.485	Oct 2009	0.584	Oct 2010	0.000		0.584	0.000	1.494	0.000
Subtotal			0.425	0.485		0.584		0.000		0.584	0.000	1.494	0.000

Remarks

Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** ZSBIR - SBIR/STTR - Aggregated from ZSBIR-SBIR/STTR	MIPR	HQ AMC, Alexandria	0.000	0.016		0.000		0.000		0.000	0.000	0.016	0.000
Subtotal			0.000	0.016		0.000		0.000		0.000	0.000	0.016	0.000

Remarks

Project Cost Totals	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
	1.582	1.302		1.821		0.000		1.821	0.000	4.705	0.000

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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	Total Prior Years Cost	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
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Remarks

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015							
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4				
** SSA - Provide Data Model Implementation Guidance	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Provide Enterprise Architecture Products and Services	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Provide Information Assurance Site Compliance Testing	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Provide Integration and Test, M&S, VV&A Certification and Accreditation	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Demonstrate Technology Transition Capabilities	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Provide CM Services for Common User Products and Services	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Provide Net-Centric Assessment and assist programs with implementation of policy	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Develop and provide CBRN Data Model implementation guidance, including reference implementations	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Architecture advisory services to support Warfighter Enterprise and Program Integrated Architectures	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Demonstrate, Verify, Test Technology Transition capabilities especially for Common Components and Services	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015							
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4				
Provide Information Assurance Certification/ Acceptance products/services, including compliance testing																																
Provide Modeling, Simulation, VV&A, Integration/Test support and interoperability demonstrations.	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Provide FISMA and J6 Interoperability certification support	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Provide CBRN Interface Standards, including reference implementations, e.g. Common CBRN Sensor Interface	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Sustain CBRN Data Model	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Sustain CCSI, including investigation, as an industry standard	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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Schedule Details

Event	Start		End	
	Quarter	Year	Quarter	Year
** SSA - Provide Data Model Implementation Guidance	1	2008	4	2015
Provide Enterprise Architecture Products and Services	3	2007	4	2015
Provide Information Assurance Site Compliance Testing	3	2006	4	2015
Provide Integration and Test, M&S, VV&A Certification and Accreditation	2	2007	4	2015
Demonstrate Technology Transition Capabilities	1	2008	4	2015
Provide CM Services for Common User Products and Services	1	2008	4	2015
Provide Net-Centric Assessment and assist programs with implementation of policy	4	2007	4	2015
Develop and provide CBRN Data Model implementation guidance, including reference implementations	1	2008	4	2015
Architecture advisory services to support Warfighter Enterprise and Program Integrated Architectures	1	2008	4	2015
Demonstrate, Verify, Test Technology Transition capabilities especially for Common Components and Services	1	2008	4	2015
Provide Information Assurance Certification/Acceptance products/services, including compliance testing	1	2008	4	2015
Provide Modeling, Simulation, VV&A, Integration/Test support and interoperability demonstrations.	1	2008	4	2015
Provide FISMA and J6 Interoperability certification support	1	2008	4	2015
Provide CBRN Interface Standards, including reference implementations, e.g. Common CBRN Sensor Interface	1	2008	4	2015

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

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Event	Start		End	
	Quarter	Year	Quarter	Year
Sustain CBRN Data Model	1	2008	4	2015
Sustain CCSI, including investigation, as an industry standard	1	2008	4	2015

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COST (\$ in Millions)	FY 2009 Actual	FY 2010 Estimate	FY 2011 Base Estimate	FY 2011 OCO Estimate	FY 2011 Total Estimate	FY 2012 Estimate	FY 2013 Estimate	FY 2014 Estimate	FY 2015 Estimate	Cost To Complete	Total Cost
TE7: <i>TEST & EVALUATION (OP SYS DEV)</i>	7.037	4.870	4.813	0.000	4.813	4.779	4.750	5.660	5.615	Continuing	Continuing
Quantity of RDT&E Articles											
A. Mission Description and Budget Item Justification											
This project provides revitalization and technology upgrades of existing instrumentation and equipment at Dugway Proving Ground (DPG), a Major Range and Test Facility Base (MRTFB), in support of their Chemical Biological test mission.											
B. Accomplishments/Planned Program (\$ in Millions)											
						FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	
1) DPG - MRTFB						1.790	0.904	0.866	0.000	0.866	
<i>FY 2009 Accomplishments:</i> Provided upgrades of the Life Sciences Test Facility instrumentation and equipment at Dugway Proving Ground (DPG), in support of their CB defense mission. This is the only U.S. facility equipped to test with aerosolized Bio-Safety Level 3 (BSL-3) agents. Upgrades and technology enhancements included: (1) Replacement of old Scanning Electron Microscopes, light microscopes, and old Aerodynamic Particle Sizers with newer Fluorescent Aerodynamic Particle Sizers. These items are replaced using a phased approach over several years; (2) Development of biological aerosols in various conditions inside the test chambers; (3) Full characterization of biological aerosols in various conditions inside the test chambers; (4) An automated aerosol dissemination systems that will vary the concentration of the aerosol cloud; (5) New methods of sampling biologics using mimetics; (6) Development of a deployable Polymerase Chain Reaction sampling system for use in the field testing of biological detection systems; (7) Continued upgrades/improvements to the Containment Aerosol Chamber (CAC) with capability to create environmental conditions with varying combinations of air											

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
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B. Accomplishments/Planned Program (\$ in Millions)

	FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
<p>temperature and relative humidity; and, (8) Continued procurement of microbiological laboratory equipment needed to utilize new Bio-Safety level 3 laboratories.</p> <p><i>FY 2010 Plans:</i> Provides upgrades of the Life Sciences Test Facility instrumentation and equipment at Dugway Proving Ground (DPG), in support of their CB defense mission. This is the only U.S. facility equipped to test with aerosolized Bio-Safety Level 3 (BSL-3) agents. Upgrades and technology enhancements include the following: (1) Replacement of a Scanning Electron Microscope and light microscopes; (2) Replacement of Aerodynamic Particle Sizers with newer Fluorescent Aerodynamic Particle Sizers; (3) Full characterization of biological aerosols in various conditions inside the test chambers; (4) An automated liquid aerosol dissemination systems that will vary concentrations of aerosols in chamber and field clouds; (5) Procure aerosol samplers for chamber and field testing; (6) Continued upgrades/improvements to the Containment Aerosol Chamber (CAC) with capability to create environmental conditions with varying combinations of air temperature and relative humidity; and, (7) Continued procurement of microbiological laboratory equipment needed to utilize new Bio-Safety level 3 laboratories.</p> <p><i>FY 2011 Base Plans:</i> Provides upgrades of the Life Sciences Test Facility instrumentation and equipment at Dugway Proving Ground (DPG), in support of their CB defense mission. This is the only U.S. facility equipped to test with aerosolized Bio-Safety Level 3 (BSL-3) agents. Upgrades and technology enhancements include the following: (1) Regular replacement of aging Aerodynamic Particle Sizers with newer Fluorescent Aerodynamic Particle Sizers; (2) Full characterization of biological aerosols in various conditions out in the field; (3) An automated dry powder dissemination system that will vary the concentration of aerosols in test chambers and in the field; (4) Procure aerosol samplers for chamber and field tests; (5) Continue to enhance genotyping system and procure genotyping analysis software; (6) Upgrade aerosol particles generation capabilities for standoff and point detector characterization;</p>					

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B. Accomplishments/Planned Program (\$ in Millions)								
				FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
and, (7) Continued procurement of microbiological laboratory equipment needed to fully utilize Bio-Safety level 3 laboratories.								
2) DPG - MRTFB <i>FY 2009 Accomplishments:</i> Provided for modernization of existing instrumentation and equipment in the major test chambers at DPG, in support of the CB defense mission. These consisted of the following: (1) the Materiel Test Facility which is a unique test chamber where real-world decontamination operations can be tested; (2) Building 4165, which houses updated chemical-surety test facilities and laboratories used for the testing of chemical protective material with chemical agents and simulants; and, (3) Building 3445, which houses two large chambers where testing of large panel decontaminants, filter systems, and Individual Protection Equipment (IPE) in a chemical environment is conducted. Modernization of instrumentation in the chambers included: (1) Continued development of a chemical aerosol generation and sampling capability; and, (2) Characterization of improved and/or articulated testing fixtures. <i>FY 2010 Plans:</i> Provides for modernization of existing instrumentation and equipment in the major test chambers at DPG, in support of the CB defense mission. These consist of the following: (1) the Materiel Test Facility which is a unique test chamber where real-world decontamination operations can be tested; (2) Building 4165, which houses updated chemical-surety test facilities and laboratories used for the testing of chemical protective material with chemical agents and simulants, and (3) Bldg 3445, which houses two large chambers where testing of large panel decontaminants, filter systems, and Individual Protection Equipment (IPE) in a chemical environment is conducted. Modernization of instrumentation in the chambers included: (1) Continued development of a chemical aerosol generation and sampling capability; (2) Characterization of improved and/or articulated testing fixtures; (3) Enhanced toxic industrial chemical detection and control capability; and (4) Initial enhancements in preparation for non traditional agent test capability.				1.980	1.020	1.059	0.000	1.059

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B. Accomplishments/Planned Program (\$ in Millions)								
				FY 2009	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total
instrumentation with current technology to include: (1) Characterization of new and upgraded test fixtures; and, (2) Upgraded control systems for small chambers. <i>FY 2010 Plans:</i> Provides for revitalization and upgrade of existing instrumentation and equipment at the Combined Chemical Test Facility at Dugway Proving Ground (DPG), in support of their CB test mission. The Combined Chemical Test Facility tests the capability of detectors, decontaminants, and protective systems to defend against toxic chemical agents. This project upgrades analytical and field instrumentation with current technology to include: (1) Characterization of new and upgraded test fixtures such as the advanced air purification fixture and novel closures fixture; (2) Upgraded control systems for small chambers such as the small item decontamination fixture; (3) Installation support for a dynamic test chamber for work with surety agents and toxic chemicals under continuously-varying conditions; (4) Characterization of upgraded real time swatch capability; and (5) Enhancement of the toxic industrial chemical test capability. <i>FY 2011 Base Plans:</i> Provides for revitalization and upgrade of existing instrumentation and equipment at the Combined Chemical Test Facility at Dugway Proving Ground (DPG), in support of their CB test mission. The Combined Chemical Test Facility tests the capability of detectors, decontaminants, and protective systems to defend against toxic chemical agents. This project upgrades analytical and field instrumentation with current technology to include: (1) Characterization of new and upgraded test fixtures; (2) Upgraded control systems for small chambers; (3) Installation support for the next-generation Chemical Biological Agent Resistance Test (CBART) capability.								
5) SBIR <i>FY 2010 Plans:</i> Small Business Innovative Research.				0.000	0.059	0.000	0.000	0.000
Accomplishments/Planned Programs Subtotals				7.037	4.870	4.813	0.000	4.813

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Exhibit R-2A, RDT&E Project Justification: PB 2011 Chemical and Biological Defense Program		DATE: February 2010
APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 7: <i>Operational Systems Development</i>	R-1 ITEM NOMENCLATURE PE 0607384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (OP SYS DEV)</i>	PROJECT TE7: <i>TEST & EVALUATION (OP SYS DEV)</i>
C. Other Program Funding Summary (\$ in Millions) N/A		
D. Acquisition Strategy T&E UPGRAD T&E Range Instrumentation/Technology Upgrades is a continuing project. It provides for technical upgrades to DPG capabilities for Chemical and Biological training and testing DoD Chemical and Biological (CB) materiel, weapons, and weapons systems from concept through production.		
E. Performance Metrics N/A		

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 7: <i>Operational Systems Development</i>	R-1 ITEM NOMENCLATURE PE 0607384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (OP SYS DEV)</i>	PROJECT TE7: <i>TEST & EVALUATION (OP SYS DEV)</i>
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Test and Evaluation (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** T&E UPGRAD - Technology Upgrades - DPG, UT	C/FP	Dugway Proving Grounds DPG, UT	13.924	4.811	Jan 2010	4.813	Jan 2011	0.000		4.813	0.000	23.548	0.000
Subtotal			13.924	4.811		4.813		0.000		4.813	0.000	23.548	0.000

Remarks

Management Services (\$ in Millions)

Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
				Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
** ZSBIR - SBIR/STTR - Aggregated from ZSBIR-SBIR/STTR	MIPR	HQ AMC, Alexandria	0.000	0.059		0.000		0.000		0.000	0.000	0.059	0.000
Subtotal			0.000	0.059		0.000		0.000		0.000	0.000	0.059	0.000

Remarks

Project Cost Totals	Total Prior Years Cost	FY 2010		FY 2011 Base		FY 2011 OCO		FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
		Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost			
Project Cost Totals	13.924	4.870		4.813		0.000		4.813	0.000	23.607	0.000

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 7: <i>Operational Systems Development</i>	R-1 ITEM NOMENCLATURE PE 0607384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (OP SYS DEV)</i>	PROJECT TE7: <i>TEST & EVALUATION (OP SYS DEV)</i>
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	Total Prior Years Cost	FY 2010	FY 2011 Base	FY 2011 OCO	FY 2011 Total	Cost To Complete	Total Cost	Target Value of Contract
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Remarks

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Exhibit R-4, RDT&E Schedule Profile: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 7: <i>Operational Systems Development</i>	R-1 ITEM NOMENCLATURE PE 0607384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (OP SYS DEV)</i>	PROJECT TE7: <i>TEST & EVALUATION (OP SYS DEV)</i>
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	FY 2009				FY 2010				FY 2011				FY 2012				FY 2013				FY 2014				FY 2015							
	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4				
** T&E UPGRAD - LSTF Instrumentation & Equip Upgrades, DPG	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Modernization of Major Test Chambers, DPG	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Enhance Instrumentation & Equip at Target S, Downwind, & Tower CB Test Grids, DPG	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■
Revitalize & Upgrade Instrumentation & Equip at Combined Chemical Test Facility, DPG	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■	■

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Exhibit R-4A, RDT&E Schedule Details: PB 2011 Chemical and Biological Defense Program **DATE:** February 2010

APPROPRIATION/BUDGET ACTIVITY 0400: <i>Research, Development, Test & Evaluation, Defense-Wide</i> BA 7: <i>Operational Systems Development</i>	R-1 ITEM NOMENCLATURE PE 0607384BP: <i>CHEMICAL/BIOLOGICAL DEFENSE (OP SYS DEV)</i>	PROJECT TE7: <i>TEST & EVALUATION (OP SYS DEV)</i>
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Schedule Details

Event	Start		End	
	Quarter	Year	Quarter	Year
** T&E UPGRAD - LSTF Instrumentation & Equip Upgrades, DPG	2	2008	2	2016
Modernization of Major Test Chambers, DPG	2	2008	2	2016
Enhance Instrumentation & Equip at Target S, Downwind, & Tower CB Test Grids, DPG	2	2008	2	2016
Revitalize & Upgrade Instrumentation & Equip at Combined Chemical Test Facility, DPG	2	2008	2	2016

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