Exhibit R-2, **RDT&E Budget Item Justification:** PB 2013 Army

APPROPRIATION/BUDGET ACTIVITY

R-1 ITEM NOMENCLATURE

2040: Research, Development, Test & Evaluation, Army

PE 0604807A: Medical Materiel/Medical Biological Defense Equipment - Eng Dev

BA 5: Development & Demonstration (SDD)

COST (\$ in Millions)	FY 2011	FY 2012	FY 2013 Base	FY 2013 OCO	FY 2013 Total	FY 2014	FY 2015	FY 2016	FY 2017	Cost To Complete	Total Cost
Total Program Element	33.262	27.132	43.395	-	43.395	46.634	47.252	48.312	49.126	Continuing	Continuing
812: MIL HIV VAC&DRUG DEV	4.297	3.860	3.232	-	3.232	4.292	4.444	4.539	4.616	Continuing	Continuing
832: Field Medical Systems Engineering Development	17.159	14.793	23.971	-	23.971	27.469	27.528	28.452	28.932	Continuing	Continuing
849: INFEC DIS DRUG/VACC ED	11.806	8.479	13.771	-	13.771	14.873	15.280	15.321	15.578	Continuing	Continuing
VS8: MEDEVAC MISSION EQUIPMENT PACKAGE (MEP) - END DEV	-	-	2.421	-	2.421	-	-	-	-	Continuing	Continuing

A. Mission Description and Budget Item Justification

This program element (PE) funds advanced development of medical materiel within the System Demonstration and Low Rate Initial Production portions of the acquisition life cycle using 6.5 funding. It supports products successfully developed in the Systems Integration portion of the Systems Development and Demonstration phases through completion of the Milestone C Decision Review. Commercially-off-the-shelf (COTS) medical products are also tested and evaluated for military use, when available. This PE primarily includes pivotal (conclusive) human clinical trials necessary for licensure by the Food and Drug Administration.

(PROJ 812) project funds military relevant human immunodeficiency virus (HIV) medical countermeasures. These funds provide for engineering and manufacturing development of candidate vaccines and drugs to permit large-scale field testing. Development focused on military unique needs effecting manning, mobilization, and deployment. Products from this project will normally transition to DoD Health Programs or OPA Funds.

(PROJ 832) this project funds the engineering and manufacturing development of medical products for enhanced combat casualty care and follow-on care, including rehabilitation. Mature commercial-off-the-shelf (COTS) medical products are also evaluated for military use. Consideration will also be given to reduce the medical sustainment footprint through smaller weight and cube volume, or equipment independence from supporting material. Products from this project will normally transition to OPA Funds.

(PROJ 849) funds development of candidate medical countermeasures for military relevant infectious diseases. These products fall between four major areas: vaccines, drugs, diagnostic kits/devices, and insect control measures to limit exposure and disease transmission. FDA approval is a mandatory obligation for all military products placed into the hands of medical providers or service members for human use. Products from this project will normally transition to DoD Health Programs or OPA funds.

(PROJ VS8) program receives products that transition from VS7 and funds effort to complete research and development for the MEDEVAC Mission Essential Packages (MEPs) to support 256 Medical Evacuation legacy helicopters. The force design will increase the number of air frames in the force from 12 to 15 aircraft for 37 MEDEVAC companies to better meet operation needs.

DATE: February 2012 Exhibit R-2, RDT&E Budget Item Justification: PB 2013 Army APPROPRIATION/BUDGET ACTIVITY R-1 ITEM NOMENCLATURE 2040: Research, Development, Test & Evaluation, Army PE 0604807A: Medical Materiel/Medical Biological Defense Equipment - Eng Dev BA 5: Development & Demonstration (SDD)

This program is managed by U.S. Army Medical Materiel Development Activity (USAMMDA) and U.S. Army Medical Materiel Agency (USAMMA) of the US Army Medical Research and Materiel Command.

B. Program Change Summary (\$ in Millions)	FY 2011	FY 2012	FY 2013 Base	FY 2013 OCO	FY 2013 Total
Previous President's Budget	34.474	27.160	41.872	-	41.872
Current President's Budget	33.262	27.132	43.395	-	43.395
Total Adjustments	-1.212	-0.028	1.523	-	1.523
 Congressional General Reductions 	-	-			
 Congressional Directed Reductions 	-	-			
 Congressional Rescissions 	-	-			
 Congressional Adds 	-	-			
 Congressional Directed Transfers 	-	-			
 Reprogrammings 	-	-			
SBIR/STTR Transfer	-0.998	-			
 Adjustments to Budget Years 	-	-	1.523	-	1.523
Other Adjustments 1	-0.214	-0.028	-	-	-

Exhibit R-2A, RDT&E Project Just	ification: PE	3 2013 Army							DATE: Febr	ruary 2012	
APPROPRIATION/BUDGET ACTIV 2040: Research, Development, Test BA 5: Development & Demonstration	& Evaluation	n, Army		PE 0604807		Materiel/Med	URE PROJECT Materiel/Medical 812: MIL HIV VAC&DRUG DEV ment - Eng Dev				
COST (\$ in Millions)	FY 2011	FY 2012	FY 2013 Base	FY 2013 OCO	FY 2013 Total	FY 2014	FY 2015	FY 2016	FY 2017	Cost To Complete	Total Cost
812: MIL HIV VAC&DRUG DEV	4.297	3.860	3.232	-	3.232	4.292	4.444	4.539	4.616	Continuing	Continuing

A. Mission Description and Budget Item Justification

Quantity of RDT&E Articles

This project funds militarily relevant human immunodeficiency virus (HIV) medical countermeasures. These funds provide for engineering and manufacturing development of candidate vaccines and drugs to permit large-scale field testing. Development is focused on militarily unique needs effecting manning, mobilization, and deployment.

The major contractor is The Henry M. Jackson Foundation for the Advancement of Military Medicine, Rockville, MD. Research efforts are coordinated with the National Institutes of Health.

B. Accomplishments/Planned Programs (\$ in Millions, Article Quantities in Each)	FY 2011	FY 2012	FY 2013
Title: Military HIV Vaccine and Drug Development	4.297	3.860	3.232
Articles:	0	0	
Description: This project provides funds for engineering and manufacturing development of candidate vaccines and drugs to permit large-scale field testing of vaccines for medical countermeasures to HIV			
FY 2011 Accomplishments: Finalized and submitted results to the Food and Drug Administration (FDA) from the expanded safety/efficacy trial in Thailand, updated/revised the clinical development plan, and continued to follow up clinical trial for HIV positive for viral load.			
FY 2012 Plans: Perform three inter-related studies to enhance our understanding of how the vaccine strategy used in the 2009 safety/efficacy trial caused vaccine recipients to be protected from infection, including intense laboratory studies using samples from the trial, and commencement of two small clinical vaccine trials to generate data and samples to define what vaccine responses to try to generate for next increment studies.			
FY 2013 Plans: Will refine vaccine administration schedule as well as clinical trial design based on data from previous clinical trials. Will adjust plan for increment 1 future efficacy trial planned to begin in late 2014.			
Accomplishments/Planned Programs Subtotals	4.297	3.860	3.232

Exhibit R-2A, RDT&E Project Justification: PB 2013 Army			DATE: February 2012
APPROPRIATION/BUDGET ACTIVITY	R-1 ITEM NOMENCLATURE	PROJECT	
2040: Research, Development, Test & Evaluation, Army	PE 0604807A: Medical Materiel/Medical	812: MIL HI	V VAC&DRUG DEV
BA 5: Development & Demonstration (SDD)	Biological Defense Equipment - Eng Dev		
0.041 D. F. II. O. (A.: 1411II.)			

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

N/A

D. Acquisition Strategy

Test and evaluate commercially developed vaccine candidates in government-managed trials.

E. Performance Metrics

Performance metrics used in the preparation of this justification material may be found in the FY 2010 Army Performance Budget Justification Book, dated May 2010.

R-1 Line #112

DATE: February 2012 Exhibit R-3, RDT&E Project Cost Analysis: PB 2013 Army APPROPRIATION/BUDGET ACTIVITY **R-1 ITEM NOMENCLATURE PROJECT** 2040: Research, Development, Test & Evaluation, Army PE 0604807A: Medical Materiel/Medical 812: MIL HIV VAC&DRUG DEV BA 5: Development & Demonstration (SDD) Biological Defense Equipment - Eng Dev FY 2013 FY 2013 FY 2013 Management Services (\$ in Millions) FY 2012 oco Base Total **Total Prior** Contract **Target** Method Performing Years Award Award Award Cost To Value of **Activity & Location Cost Category Item** & Type Cost Cost Date Cost Date Cost Date Complete **Total Cost** Contract Cost No product/contract costs Various:Various 1.124 0.215 0.299 0.299 Continuing Continuing 0.000 Various greater than \$1M individually Subtotal 1.124 0.215 0.299 0.299 0.000 **FY 2013** FY 2013 FY 2013 **Product Development (\$ in Millions)** FY 2012 Base oco Total **Total Prior** Contract **Target** Method Performing Years Award Award Award **Cost To** Value of **Cost Category Item** & Type **Activity & Location** Cost Cost Date Cost Date Cost Date Cost Complete **Total Cost** Contract Henry M. Jackson **Product Development** Various 2.655 2.145 2.145 Continuing 27.742 Continuing Continuing Foundation,:Various Subtotal 27.742 2.655 2.145 2.145 FY 2013 FY 2013 FY 2013 Support (\$ in Millions) FY 2012 Base oco Total Contract **Total Prior Target** Method Performing Years Award Award Award Cost To Value of **Cost Category Item** & Type **Activity & Location** Cost Cost Date Cost Date Cost Date Cost Complete **Total Cost** Contract No product/contract costs 0.046 0.031 Continuing Various Various:Various 0.580 0.031 Continuing 0.000 greater than \$1M individually Subtotal 0.580 0.046 0.031 0.031 0.000 FY 2013 FY 2013 FY 2013 Test and Evaluation (\$ in Millions) FY 2012 Base oco Total Contract **Total Prior Target** Method Performing Cost To Value of Years Award Award Award **Cost Category Item** & Type **Activity & Location** Cost Cost Date Cost Date Cost Date Cost Complete **Total Cost** Contract Henry M. Jackson **Product Development** Various 23.446 0.944 0.757 0.757 Continuing Continuing Continuing Foundation.: Various 23.446 0.944 0.757 0.757 Subtotal

Exhibit R-3, RDT&E Project Cost Analysis: PB 2013 Army			DATE: February 2012
APPROPRIATION/BUDGET ACTIVITY	R-1 ITEM NOMENCLATURE	PROJECT	
2040: Research, Development, Test & Evaluation, Army	PE 0604807A: Medical Materiel/Medical	812: MIL HI	IV VAC&DRUG DEV
BA 5: Development & Demonstration (SDD)	Biological Defense Equipment - Eng Dev		

	Total Prior Years		FY 2013	FY 2013	FY 2013	Cost To		Target Value of
	Cost	FY 2012	Base	oco	Total	Complete	Total Cost	Contract
Project Cost Total	ls 52.892	3.860	3.232	-	3.232			

Remarks

Exhibit R-4, RDT&E Schedule Profile: PB 2013 Army

APPROPRIATION/BUDGET ACTIVITY

2040: Research, Development, Test & Evaluation, Army
BA 5: Development & Demonstration (SDD)

R-1 ITEM NOMENCLATURE
PE 0604807A: Medical Materiel/Medical
Biological Defense Equipment - Eng Dev

DATE: February 2012

PROJECT
812: MIL HIV VAC&DRUG DEV

		FY	2011	1		FY 2012			FY 2013			FY 2	2014	FY 2015			,	FY 2016			,	FY 2017			,			
	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
Phase 2 study of Vaccine candidates				·	·					,									·									
Initiate Phase 3 Study of Vaccine candidates																												
Increment 1 Efficacy Trial																												

Exhibit R-4A, RDT&E Schedule Details: PB 2013 Army

APPROPRIATION/BUDGET ACTIVITY

2040: Research, Development, Test & Evaluation, Army

BA 5: Development & Demonstration (SDD)

PATE: February 2012

R-1 ITEM NOMENCLATURE

PE 0604807A: Medical Materiel/Medical
Biological Defense Equipment - Eng Dev

PROJECT

812: MIL HIV VAC&DRUG DEV

Schedule Details

	St	art	E	nd
Events	Quarter	Year	Quarter	Year
Phase 2 study of Vaccine candidates	1	2014	2	2014
Initiate Phase 3 Study of Vaccine candidates	1	2015	1	2015
Increment 1 Efficacy Trial	4	2014	2	2015

Exhibit R-2A, RDT&E Project Jus	tification: PE	3 2013 Army							DATE : Febr	uary 2012	
APPROPRIATION/BUDGET ACTIV 2040: Research, Development, Tes BA 5: Development & Demonstration	t & Evaluation	n, Army			I OMENCLA 7A: <i>Medical</i> Defense Equi	Materiel/Med		PROJECT 832: Field M Developme	Medical Syste	ering	
COST (\$ in Millions)	FY 2011	FY 2012	FY 2013 Base	FY 2013 OCO	FY 2013 Total	FY 2014	FY 2015	FY 2016	FY 2017	Cost To Complete	Total Cost
832: Field Medical Systems Engineering Development	17.159	14.793	23.971	-	23.971	27.469	27.528	28.452	28.932	Continuing	Continuing
Quantity of RDT&E Articles											

A. Mission Description and Budget Item Justification

This project funds the engineering and manufacturing development of medical products for enhanced combat casualty care and follow-on care, including rehabilitation. This project funds pivotal (conclusive) human clinical trials or mechanical engineering evaluations for efficacy of devices or biologics (products derived from living organisms) to fulfill unique military requirements. Mature commercial-off-the-shelf (COTS) medical products are also evaluated for military use. Consideration will also be given to reducing the medical sustainment footprint through smaller weight and cube volume, or equipment independence from supporting materiel. This work is frequently completed through a laboratory/contractor team with the contractor obtaining the U.S. Food and Drug Administration (FDA) licensure for sale of the product.

Major contractors/intra-governmental agencies include: IGR Enterprises, Inc.; Army Medical Department Board Test Center; SeQual Technologies, Inc.; Enginivity, Inc.; Ultrasound Diagnostics, Inc.; HemCon Medical Technologies, Inc.; Cerdak Ltd; Hemerus Medical, LLC; Fast Track Drugs & Biologics, LLC; Integrated Medical Systems, Inc; the National Institutes of Health National Heart, Lung and Blood Institute (NHLBI), and the U.S. Army Aeromedical Research Laboratory Walter Reed Army Institute of Research (WRAIR) and Institute of Surgical Research (ISR) for user evaluation. Other military agencies include Program Executive Office (PEO) Soldier, PEO Combat Service Support (CSS), and Naval Undersea Warfare Center.

B. Accomplishments/Planned Programs (\$ in Millions, Article Quantities in Each)	FY 2011	FY 2012	FY 2013
Title: Medical Materiel/Medical Biological Defense Equipment- Engineering Development FY 2010 and 2011	17.159	_	-
Articles:	0		
Description: This project funds in FY 2010 and 2011 the engineering and manufacturing development of medical products for enhanced combat casualty care and follow-on care, including rehabilitation.			
FY 2011 Accomplishments:			
In FY 2011, Freeze-dried Plasma: start enrollment in the multi-center limited human safety/efficacy clinical trial and evaluate			
stability profile of the product. Red Blood Cell Extended Life Program: complete the human safety/efficacy study, conduct data			
analysis, and prepare and submit report to the FDA. Platelet Derived Hemostatic Agent (PDHA): transition from project 836 to			
begin enrollment/follow-up in the human safety/efficacy trial. Intranasal Ketamine (low dose pain management via nasal spray):			
transition from project 836 and conduct pre-clinical trial activities for human drug use in aeromedical evacuation and burn patients.			
Hypertonic Saline Dextran (treatment of shock due to blood loss): support human expanded safety/efficacy trial for treatment of			
shock. Continue market analysis and surveys for candidate resuscitative fluids for treatment of shock. Enhanced SAVe: Conduct clinical user evaluation and operational testing. Remote Diagnostic Access: Conduct Prototype field testing and refinement for			
clinical user evaluation and operational testing. Remote Diagnostic Access. Conduct Prototype field testing and reinfernent for			

Exhibit R-2A, RDT&E Project Justification: PB 2013 Army			DATE: Fe	bruary 2012	
APPROPRIATION/BUDGET ACTIVITY 2040: Research, Development, Test & Evaluation, Army BA 5: Development & Demonstration (SDD)	R-1 ITEM NOMENCLATURE PE 0604807A: Medical Materiel/Medical Biological Defense Equipment - Eng Dev	PROJECT 832: Field Developm	Medical Sys	stems Engine	ering
B. Accomplishments/Planned Programs (\$ in Millions, Article	Quantities in Each)		FY 2011	FY 2012	FY 2013
MOC Sensor Modual Development phase of this device. Treatmet Variants: Transition from Project 836. Transition to production for HBCT for Medical Mission Package for Treatment Variant: Transitesting. Shelter for PM, HBCT Treatment Variant: Transition from I Shock and Vibration Isolation System for Patient Litters in Ground 836. Conduct Shock and Vibration Isolation System for patient litt tourniquet update: Conduct study to update medical set and down Sensitive Medical Materiel: Transition from Project 836. Conduct	use on PM, HBCT Treatment Variant. Refrigerator for tion from Project 836. Conduct environmental and operation 836. Evaluate alternatives in operational set and Air Medical Evacuation Vehicles: Transition from ers - air worthiness and ground vehicle testing. One-select product. Passive Cold Chain Capability for Testing 1981.	or PM, perational ting. n Project handed			
Title: Field Medical Systems Engineering Development FY 2012 F	PM Medical Devices	Articles:	-	6.019 0	0.20
Description: This project funds in FY 2012 the engineering and m combat casualty care managed by PM Medical Devices.	nanufacturing development of medical products for er				
FY 2012 Plans: The Burn Resuscitation Decision Device: Will conduct final enviror submission. Wireless Medical Monitoring will transition from Congresion project and will undergo a Milestone C review as well as prototype line into 832 6.5 funding line. Will finalize results of clinical and op and undergo Milestone C review in 4Q.	ressional Special Interest (CSI) project to an Army Co e field testing. Plasma Knife: Will transition from 836	ore funded 6.4 funding			
FY 2013 Plans: Enhanced SAVE: No further R&D funding required as this is now Preparing documentation for CPD and MS B/C. No further R&D re 2013. MS B/C expected 2Q FY 2013. Total Intravenus Anesthsia (Wireless Medical Monitoring: no requirement and no RDTE planne a COTS product. Noise-Immune Stethoscope: Will finalize and co Oxygen Generator (15 LPM) System: 15LPM draft CDD complete 2012. Will continue development with a target to field in FY 2015. planned for testing of the device. Will begin design and developm 2013. One-Handed Tourniquet update: No RDTE efforts in FY 2015.	equired as this product will transition to procurement in (TIVA): This product is transitioning to tech watch in ed. Plasma Knife: No further R&D funding required as and a MS C review in FY 2012; no RDTE efforts for d and a request for proposals (RFP) award is expected. Replacement for the M-138 Steam Sterilizer: FY 20 tent of system in FY 2012 and continue development.	n FY FY 2012. s this is now or FY 2013. ed in March 13 funding			
Title: Field Medical Systems Engineering Development FY 2012 F		Articles:	-	5.034 0	17.59

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Exhibit R-2A, RDT&E Project Justification: PB 2013 Army			DATE: Fe	bruary 2012	
APPROPRIATION/BUDGET ACTIVITY 2040: Research, Development, Test & Evaluation, Army BA 5: Development & Demonstration (SDD)	R-1 ITEM NOMENCLATURE PE 0604807A: Medical Materiel/Medical Biological Defense Equipment - Eng Dev	PROJECT 832: Field I Developme	•	stems Engine	ering
B. Accomplishments/Planned Programs (\$ in Millions, Art	ticle Quantities in Each)		FY 2011	FY 2012	FY 2013
Description: Funding is provided for engineering and manufal Pharmaceuticals for enhanced combat casualty care and follows:		1			
FY 2012 Plans: Freeze-Dried Plasma: Begin enrollment in a multi-center limite manufacturing & laboratory capabilities to meet Food and Dru the product. Cryopreserved Platelets (CPP) (formerly Platelet in a human safety/efficacy clinical trial.	ug Administration requirements; and will evaluate stability p	orofile of			
FY 2013 Plans: Blood Pathogen Reduction/Inactivation transitioned to advance with Defense Health Program RDT&E funding; transitioned to avoid delays. Freeze-Dried Plasma: Finalize Phase 3 test plan and continue development of commercially sustainable currer Accelerated fielding of a FDA-approved Freeze-Dried Plasma the Blast Injury Task Force. Cryopreserved Platelets: Continuprocesses in support of U.S. Food and Drug Administration lice Phase 3 Pivotal clinical trial is required by the U.S. Food and	Freeze-dried Plasma program to maintain current schedun and protocols, recruit test sites for Phase 3 Pivotal clinicant Good Manufacturing Practices compliant manufacturing was validated in the June 2011 Army Surgeon General's use validation of current Good Manufacturing Practices marcensure. Develop Phase 3 clinical testing network and processing process.	le and al trial, capability. Report by nufacturing			
Title: Field Medical Systems Engineering Development FY 20	012 PM Integrated Clinical Systems (ICS)	Articles:	-	0.737 0	-
Description: This project funds in FY 2012 the engineering a ICS for enhanced combat casualty care and follow-on care, in				0	
FY 2012 Plans: Will conduct final testing of Phase IV of Remote Acess Device to serve as a standardized compliance wrapper for all vendors traverse between the .com and .mil networks. The Milestone I RDA project in March 2011.	s, medical technology, and even IT management products	that may			
Title: Field Medical Systems Engineering Development FY 20	012 PM Medical Support Systems	Articles:	-	3.003	6.172
Description: This project funds in FY 12 the engineering and Medical Support Systems for enhanced combat casualty care				U	

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Exhibit R-2A, RDT&E Project Justification: PB 2013 Army		DATE: February 2012
APPROPRIATION/BUDGET ACTIVITY	R-1 ITEM NOMENCLATURE	PROJECT
2040: Research, Development, Test & Evaluation, Army	PE 0604807A: Medical Materiel/Medical	832: Field Medical Systems Engineering
BA 5: Development & Demonstration (SDD)	Biological Defense Equipment - Eng Dev	Development

B. Accomplishments/Planned Programs (\$ in Millions, Article Quantities in Each)	FY 2011	FY 2012	FY 2013
FY 2012 Plans: Collaborate with PM HBCT on medical equipment for the future treatment vehicle variant. Test and evaluate the shock and litter isolation system for potential addition to Ground Ambulance and Air Ambulance Medical Equipment Sets (MES). Transition to Force Provider by fielding a Force Provider CSH. Fully test the new 2-sided ISO shelter under the Joint shelter Program and finalize acquisition strategy in coordination with Program Manager's Force Provider program. Continue to develop Medical Evacuation Vehicles in coordination with Program Executive Office Combat Service Support vehicle developers.			
Will transition from 836 and collaborate with PM HBCT on final integration/operational testing of the treatment table and blood refrigerator in the future treatment vehicle variant. As part of the medical equipment sets, will transition cold chain technology, trauma tiered medical bag, water/waste water management system, and quad fold litter from 836 and complete operational evaluation. Will continue modernization of medical equipment sets for preventive medicine, air and ground medical evacuation, and fresh water/waste water combat support hospital support. Will transition ISO panel from 836 and complete operational testing. Will transition from 836 and complete final operational evaluation of Force Provider CSH. Will complete operational/technical testing of Future Medical Shelter System (hard-wall 1-sided and 2-sided shelters) for a material procurement decision. Will continue collaboration with PEO Combat Service Support on finalization of MRAP medical vehicle evacuation platforms including a redesign of the Casualty Evacuation (CASEVAC) medical equipment set.			
Accomplishments/Planned Programs Subtotals	17.159	14.793	23.971

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

N/A

D. Acquisition Strategy

Develop in-house or industrial prototypes in government-managed programs to meet military and regulatory requirements for production and fielding.

E. Performance Metrics

Performance metrics used in the preparation of this justification material may be found in the FY 2010 Army Performance Budget Justification Book, dated May 2010.

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Exhibit R-3, RDT&E Project Cost Analysis: PB 2013 Army

APPROPRIATION/BUDGET ACTIVITY

2040: Research, Development, Test & Evaluation, Army

BA 5: Development & Demonstration (SDD)

R-1 ITEM NOMENCLATURE

PE 0604807A: Medical Materiel/Medical

Biological Defense Equipment - Eng Dev

DATE: February 2012

PROJECT

832: Field Medical Systems Engineering

Development

27 (0. 2 0 1 0 1 0 1 1 0 1 1 0 1 1 0 1	0111011011011011	(322)			09.00. 20.	5.7.00 <u>= qu.,p</u>		9 201	20.0	<i>-</i>			
Management Services	s (\$ in Millic	ons)		FY 2	2012	FY 2 Ba			2013 CO	FY 2013 Total			
Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost	Cost To	Total Cost	Target Value of Contract
No product/contract costs greater than \$1Million individually	Various	Various:Various	22.060	0.418		2.577		-		2.577	Continuing	Continuing	Continuing
	<u>'</u>	Subtotal	22.060	0.418		2.577		-		2.577			
Product Development	(\$ in Millio	ons)		FY 2	2012	FY 2 Ba			2013 CO	FY 2013 Total			
Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost	Cost To Complete	Total Cost	Target Value of Contract
		HemCon Medical											

Troduct Development (\$\psi\$ in willions)			FY 2	2012	Ва	se	0	CO	Total				
Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost	Cost To Complete	Total Cost	Target Value of Contract
Freeze-dried Human Plasma	Various	HemCon Medical Technologies, Inc,:Tigard OR	17.525	5.796		6.000		-		6.000	Continuing	Continuing	Continuing
Hypertonic Saline Dextran	Various	National Institutes of Health, National Heart, Lung and Blood Institute (NHLBI):Various	15.100	-		-		-		-	Continuing	Continuing	Continuing
Extended Life Red Blood Cell Product	Various	Hemerus Medical, LLC,:Various	3.140	-		-		-		-	Continuing	Continuing	Continuing
No product/contract costs greater than \$1M individually	Various	Various:Various	3.260	-		0.250		-		0.250	Continuing	Continuing	Continuing
Cryopreserved Platelets	Various	Multiple DoD activities and Dartmouth Hitchcock Med Ctr:North Potomac, MD	-	7.757		9.108		-		9.108	Continuing	Continuing	Continuing
		Subtotal	39.025	13.553		15.358		-		15.358			

Exhibit R-3, RDT&E Project Cost Analysis: PB 2013 Army

APPROPRIATION/BUDGET ACTIVITY

2040: Research, Development, Test & Evaluation, Army

BA 5: Development & Demonstration (SDD)

R-1 ITEM NOMENCLATURE

PE 0604807A: Medical Materiel/Medical Biological Defense Equipment - Eng Dev **PROJECT**

832: Field Medical Systems Engineering

DATE: February 2012

Development

Support (\$ in Millions))			FY 2	2012	FY 2 Ba			2013 CO	FY 2013 Total			
Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost	Cost To Complete	Total Cost	Target Value of Contract
Regulatory Support	Various	Clinical Research Management,Inc,.:Variou	5.557	-		-		-		-	Continuing	Continuing	Continuing
No product/contract costs greater then \$1 Million individually	Various	Various:Various	4.328	0.418		1.108		-		1.108	Continuing	Continuing	Continuing
		Subtotal	9.885	0.418		1.108		-		1.108			
Took and Fredrick (A						FY 2	013	FY:	2013	FY 2013			

Test and Evaluation (\$	in Millions)		FY 2	012	FY 2 Ba	2013 se		2013 CO	FY 2013 Total			
Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost	Cost To Complete	Total Cost	Target Value of Contract
No product/contract costs greater than \$1 Million individually	Various	Various:Various	7.292	0.404		4.928		-		4.928	Continuing	Continuing	Continuing
		Subtotal	7.292	0.404		4.928		-		4.928			

_								
-	Total Prior							Target
	Years		FY 2013	FY 2013	FY 2013	Cost To		Value of
	Cost	FY 2012	Base	oco	Total	Complete	Total Cost	Contract
Project Cost Totals	78 262	14 793	23 971	_	23 971			

Remarks

Exhibit R-4, RDT&E Schedule Profile: PB 2013 Army					DATE: Februar	ry 2012	
APPROPRIATION/BUDGET ACTIVITY 2040: Research, Development, Test & Evaluation, Army BA 5: Development & Demonstration (SDD)	PE 0604807	OMENCLATUR 7A: Medical Mat Defense Equipme	eriel/Medical	PROJECT 832: Field I Developme	Medical Systems	s Engineering	
	-V 2042	EV 2042	EV 2044	EV 2045	EV 2040	EV 2047	—— 1

		FY 2011		FY 2011 FY 2012			FY 2013			FY 2014		FY 2015				FY 2016				FY 2017							
	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
Total Intravenous Anesthesia (TIVA) (MS-C)		·	,								·	,							,,								
Burn Resuscitation Decision Device (MS-C)																											
Wireless Medical Monitoring (MS-C)																											
Plasma Knife (MS-C)																											

Exhibit R-4A, RDT&E Schedule Details: PB 2013 Army			DATE: February 2012
APPROPRIATION/BUDGET ACTIVITY	R-1 ITEM NOMENCLATURE	PROJECT	
2040: Research, Development, Test & Evaluation, Army	PE 0604807A: Medical Materiel/Medical	832: Field N	Medical Systems Engineering
BA 5: Development & Demonstration (SDD)	Biological Defense Equipment - Eng Dev	Developme	nt

Schedule Details

	Sta	art	Eı	nd
Events	Quarter	Year	Quarter	Year
Total Intravenous Anesthesia (TIVA) (MS-C)	1	2013	1	2013
Burn Resuscitation Decision Device (MS-C)	2	2012	2	2012
Wireless Medical Monitoring (MS-C)	4	2012	4	2012
Plasma Knife (MS-C)	2	2012	2	2012

Exhibit R-2A, RDT&E Project Just	ification: PE	3 2013 Army							DATE: Febr	uary 2012				
APPROPRIATION/BUDGET ACTIV 2040: Research, Development, Test BA 5: Development & Demonstration	& Evaluation	n, Army		PE 060480	IOMENCLA [*] 7A: Medical Defense Equi	Materiel/Med		PROJECT 849: INFEC DIS DRUG/VACC ED						
COST (\$ in Millions)	FY 2011	FY 2012	FY 2013 Base	FY 2013 OCO	FY 2013 Total	FY 2014	FY 2015	FY 2016	FY 2017	Cost To Complete	Total Cost			
849: INFEC DIS DRUG/VACC ED	11.806	8.479	13.771	-	13.771	14.873	15.280	15.321	15.578	Continuing	Continuing			
Quantity of RDT&E Articles														

A. Mission Description and Budget Item Justification

This project funds development of candidate medical countermeasures for militarily relevant infectious diseases. These products fall within four major areas: vaccines, drugs, diagnostic kits/devices, and insect control measures to limit exposure and disease transmission. It funds research that supports conclusive human clinical trials for large-scale human efficacy testing, expanded human safety clinical trials, long-term animal studies, and related manufacturing tests. This work, which is jointly performed by military laboratories, civilian contracted pharmaceutical firms and foreign research partners, is directed toward the prevention of disease, early diagnosis, and speeding recovery once diagnosed. Medical products approved for human use must successfully complete a series of clinical trials that are required and regulated by the U.S. Food and Drug Administration (FDA). FDA approval is a mandatory obligation for all military products placed into the hands of medical providers or service members for human use. Development priority is based upon four major factors: (1) the extent of the disease within the Combatant Commands' theater of operations, (2) the clinical severity of the disease, (3) the technical maturity of the proposed solution, and (4) the affordability of the solution (development, production, and sustainment). Malaria, dysentery, hepatitis, and dengue diseases (a severe debilitating disease transmitted by mosquitoes), which are found in Africa Command, Central Command, European Command, Southern Command, and Pacific Command areas are at the top of the infectious diseases requirements list.

B. Accomplishments/Planned Programs (\$ in Millions, Article Quantities in Each)	FY 2011	FY 2012	FY 2013
Title: FY 2010 Malaria Drug/ Vaccine Engineering Development	4.147	-	-
Articles:	0		
Description: This project funds development of candidate medical countermeasures for militarily relevant infectious diseases.			
FY 2011 Accomplishments:			
Conducted clinical trials, developmental testing, and reviews of malarial/antimalarial vaccines, drugs, diagnostics and insect			
repellents. For Tafenoquine, completed pre-trial activities for a large-scale safety/efficacy human clinical trial (treatment			
indication) in a malaria endemic country.			
Title: FY 2010 Infectious Disease Drug and Vaccine Engineering Development	7.659	8.479	13.771
Articles:	0	0	
Description: FY 2010 funding for research and development efforts described below.			
FY 2011 Accomplishments:			
Analyzed the data for Topical Antileishmanial Cream, and prepared the final report on the Tunisia large scale human safety/			
efficacy trial, continued pretrial activities for enrollment in the second large scale (> 300 subjects) human safety/efficacy trial in			
Central/South America. Analyzed the data and prepared the final report for the three small scale (<35 subjects) pharmacokinetic			

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Exhibit R-2A, RDT&E Project Justification: PB 2013 Army		DATE: February 2012	
APPROPRIATION/BUDGET ACTIVITY 2040: Research, Development, Test & Evaluation, Army	R-1 ITEM NOMENCLATURE PE 0604807A: Medical Materiel/Medical	PROJECT 849: INFEC	DIS DRUG/VACC ED
BA 5: Development & Demonstration (SDD)	Biological Defense Equipment - Eng Dev	040. 777 20	, bio bitos, vitos eb

B. Accomplishments/Planned Programs (\$ in Millions, Article Quantities in Each)	FY 2011	FY 2012	FY 2013
(drug metabolism) human trials in France, Peru and Panama. Continued a human treatment protocol in the U.S For Pentostam.			
Completed the study report for DTV vaccine on the expanded human safety and immunity trial in Puerto Rico and began the expanded safety and efficacy trial in children in Thailand. Continued maintenance of study blood specimens for Hepatitis E			
vaccine kept under cold storage and monitored industry partner's Hepatitis E vaccine development effort.			
FY 2012 Plans: Conduct Clinical trials, developmental testing, and reviews of malarial/antimalarial vaccines, drugs, diagnostics and insect			
repellents. Down-select from candidate anti-malaria drugs (e.g. Tafenoquine and other drugs) to prepare for clinical trial activities			
for a safety/efficacy human clinical trial (treatment indication) in a malaria endemic country. Conduct clinical trials, developmental testing, and appropriate reviews of grouped vaccines, drugs, and diagnostics (Leishmaniasis (a skin-based disease caused by a			
parasite and transmitted by sand flies), Dengue (a severe debilitating disease caused by a virus and transmitted by a mosquito),			
and other viral diseases: For Topical Antileishmanial Cream, begin the clinical study report on the Tunisia large scale human safety/efficacy trial, begin enrollment efforts in the second large scale (> 300 subjects) human safety/efficacy trial in Central/South			
America. For Dengue Tetravalent Vaccine (DTV), begin study close out activities for expanded safety and efficacy trial in children			
in Thailand; begin a new DTV large scale (> 300 subjects) human trial in Thailand and Philippines to expand safety data and			
demonstrate efficacy in children/adults.			
FY 2013 Plans: Review and analyze data from the on-going Adult Indication study begun in FY 2012 with industry partner Sanofi Pasteur			
and determine a Go/No Go Decision on continued product development for the Dengue Tetravalent Vaccine. Phase 3 clinical			
effectiveness studies are on-going with industry partner Sanofi Pasteur for the Dengue Tetravalent Vaccine, as well as Phase 3 studies for traveler/military indication. Complete preparation prior to initiating Phase 3 Pivotal clinical trial for Malaria Prophylaxis			
Drug. For Topical Antileishmanial Cream, complete Phase 2 safety and effectiveness New World clinical trial analysis and			
complete Phase 3 New World Pivotal clinical trial, and begin New World Treatment Protocol for Phase 3 site(s). The enteric JBAIDS assay transitions to advanced development and clinical trial planning begins. The Dengue Rapid Diagnostic Device			
(DRDD) (Hand Held Infectious Disease Diagnostics) transitions to advanced development and will be evaluated in a multi-			
site clinical performance study. Leishmania Rapid Diagnostic Device (LRDD) will continue the new world clinical trial started in			
FY 2012. The Leishmania Skin Test project will complete FDA approval and transition to procurement. The Antimalarial Drug, Artesunate Intravenous will transition from 808 and conduct MS C review.			
Accomplishments/Planned Programs Subtotals	11.806	8.479	13.771

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

N/A

Exhibit R-2A, RDT&E Project Justification: PB 2013 Army		DATE: February 2012
APPROPRIATION/BUDGET ACTIVITY 2040: Research, Development, Test & Evaluation, Army BA 5: Development & Demonstration (SDD)	R-1 ITEM NOMENCLATURE PE 0604807A: Medical Materiel/Medical Biological Defense Equipment - Eng Dev	PROJECT 849: INFEC DIS DRUG/VACC ED
 <u>D. Acquisition Strategy</u> Test and evaluate in-house and commercially developed produ registration. 	cts in government-managed trials to meet FDA requir	ements and Environmental Protection Agency
E. Performance Metrics		
Performance metrics used in the preparation of this justification	material may be found in the FY 2010 Army Performa	ance Budget Justification Book, dated May 2010.

PE 0604807A: Medical Materiel/Medical Biological Defense Equipm... Army

DATE: February 2012 Exhibit R-3, RDT&E Project Cost Analysis: PB 2013 Army APPROPRIATION/BUDGET ACTIVITY **R-1 ITEM NOMENCLATURE PROJECT** 2040: Research, Development, Test & Evaluation, Army PE 0604807A: Medical Materiel/Medical 849: INFEC DIS DRUG/VACC ED BA 5: Development & Demonstration (SDD) Biological Defense Equipment - Eng Dev FY 2013 FY 2013 FY 2013 Management Services (\$ in Millions) oco FY 2012 Base Total **Total Prior** Contract Target Method Performing Years Award Award Award Cost To Value of **Activity & Location** Cost Complete **Cost Category Item** Cost Date Cost Date Date **Total Cost** Contract & Type Cost Cost No product/contract costs 12.558 1.931 2.172 2.172 Continuing Various Various: Various Continuing Continuing greater than \$1M individually Subtotal 12.558 1.931 2.172 2.172 **FY 2013** FY 2013 FY 2013 **Product Development (\$ in Millions)** FY 2012 Base oco Total **Total Prior** Contract **Target** Method Performing Years Award Award Award **Cost To** Value of **Cost Category Item** & Type **Activity & Location** Cost Cost Date Cost Date Cost Date Cost Complete **Total Cost** Contract No product/contract costs Various:Various 4.034 Continuing Various 21.423 3.171 4.034 Continuing Continuing greater than \$1M individually Topical Antileishmanial Drug **TBD** TBD:TBD 2.400 2.400 0.000 2.400 0.000 Subtotal 21.423 3.171 6 434 6 434 FY 2013 **FY 2013** FY 2013 Support (\$ in Millions) FY 2012 Base oco Total **Total Prior** Contract Target Method Years Cost To Value of Performing Award Award Award **Cost Category Item** & Type **Activity & Location** Cost Cost Date Cost Date Complete **Total Cost** Contract Cost Date Cost No product/contract costs Various Various Various 10.951 0.992 2 620 2.620 Continuina Continuina Continuina greater than \$1M individually Subtotal 10.951 0.992 2.620 2.620 _ FY 2013 FY 2013 FY 2013 Test and Evaluation (\$ in Millions) FY 2012 Base oco Total **Total Prior** Contract Target Method **Cost To** Value of Performing Years Award Award Award **Cost Category Item** & Type **Activity & Location** Cost Cost Date Cost Date Cost Date Cost Complete **Total Cost** Contract No product/contract costs 2.385 2.545 2.545 31.778 Continuina Continuina Continuina Various Various:Various greater than \$1M individually 2.385 2.545 2.545 Subtotal 31.778

Exhibit R-3, RDT&E Project Cost Analysis: PB 2013 Army	DATE: February 2012		
APPROPRIATION/BUDGET ACTIVITY	R-1 ITEM NOMENCLATURE	PROJECT	
2040: Research, Development, Test & Evaluation, Army	PE 0604807A: Medical Materiel/Medical	849: <i>INFEC</i>	DIS DRUG/VACC ED
BA 5: Development & Demonstration (SDD)	Biological Defense Equipment - Eng Dev		

, ,								
	Total Prior							Target
	Years		FY	2013 FY	2013 FY 2013	Cost To		Value of
	Cost	FY 2	2012 Ba	ase O	CO Total	Complete	Total Cost	Contract
Project Cost Totals	76.710	8.479	13.771	-	13.771			

Remarks

Exhibit R-4, RDT&E Schedule Profile: PB 2013 Army

APPROPRIATION/BUDGET ACTIVITY

2040: Research, Development, Test & Evaluation, Army
BA 5: Development & Demonstration (SDD)

PATE: February 2012

R-1 ITEM NOMENCLATURE
PE 0604807A: Medical Materiel/Medical
Biological Defense Equipment - Eng Dev

PROJECT
849: INFEC DIS DRUG/VACC ED

		FY 2	2011			F١	20 1	12		FY 2013 FY 2014						FY 2015 F				FY	201		FY 2017						
	1	2	3	4	1	1	2 3	3 4	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
Dengue Tetravalent Vaccine (DTV) Critical Design Review (CDR)																													
DTV Phase 3 Adult Indication Studies																													
DTV Adult Indication Decision																													
DTV Milestone C (MS-C)																													
DTV Biologic Licensing Application (BLA) Submission																													
Malaria Prophylaxis Phase 3 Pivotal Clinical Trial																													
Malaria Prophylaxis (MS-C)																													
Paromomycin/Gentamicin Topical Antileishmanial Cream (TLC)																													
Paromomycin/Gentamicin TLC Phase 3 Clinical Trials																													
Paromomycin/Gentamicin TLC Milestone C (MS-C)																													
Paromomycin/Gentamicin TLC New Drug Application (NDA)																													
Paromomycin/Gentamicin TLC FDA Approval																													
Paromomycin/Gentamicin TLC																													
Leishmania Rapid Diagnostic Device MS-C																													
Leishmania Rapid Diagnostic Device FDA Approval																													
Leishmania Rapid Diagnostic Device Fielding/ Delivery																													

DATE: February 2012 Exhibit R-4A, RDT&E Schedule Details: PB 2013 Army

APPROPRIATION/BUDGET ACTIVITY R-1 ITEM NOMENCLATURE **PROJECT**

2040: Research, Development, Test & Evaluation, Army PE 0604807A: Medical Materiel/Medical 849: INFEC DIS DRUG/VACC ED

BA 5: Development & Demonstration (SDD) Biological Defense Equipment - Eng Dev

Schedule Details

	Sta	Start		ıd
Events	Quarter	Year	Quarter	Year
Dengue Tetravalent Vaccine (DTV) Critical Design Review (CDR)	4	2011	4	2011
DTV Phase 3 Adult Indication Studies	2	2012	2	2015
DTV Adult Indication Decision	4	2013	4	2013
DTV Milestone C (MS-C)	4	2016	4	2016
DTV Biologic Licensing Application (BLA) Submission	4	2017	4	2017
Malaria Prophylaxis Phase 3 Pivotal Clinical Trial	4	2013	4	2015
Malaria Prophylaxis (MS-C)	1	2017	1	2017
Paromomycin/Gentamicin Topical Antileishmanial Cream (TLC)	1	2012	1	2014
Paromomycin/Gentamicin TLC Phase 3 Clinical Trials	3	2011	3	2013
Paromomycin/Gentamicin TLC Milestone C (MS-C)	1	2014	1	2014
Paromomycin/Gentamicin TLC New Drug Application (NDA)	4	2014	4	2014
Paromomycin/Gentamicin TLC FDA Approval	4	2015	4	2015
Paromomycin/Gentamicin TLC	1	2016	4	2017
Leishmania Rapid Diagnostic Device MS-C	1	2014	1	2014
Leishmania Rapid Diagnostic Device FDA Approval	2	2014	2	2014
Leishmania Rapid Diagnostic Device Fielding/Delivery	4	2014	4	2015

Exhibit R-2A, RDT&E Project Just	ification: PE	3 2013 Army	,						DATE: Feb	ruary 2012			
APPROPRIATION/BUDGET ACTIV 2040: Research, Development, Test BA 5: Development & Demonstration	& Evaluation	n, Army			IOMENCLA 7A: Medical Pefense Equi	Materiel/Med	PROJECT VS8: MEDEVAC MISSION EQUIPMENT PACKAGE (MEP) - END DEV						
COST (\$ in Millions)	FY 2011	FY 2012	FY 2013 Base	FY 2013 OCO	FY 2013 Total	FY 2014	FY 2015	FY 2016	FY 2017	Cost To Complete	Total Cost		
VS8: MEDEVAC MISSION EQUIPMENT PACKAGE (MEP) - END DEV	-	-	2.421	-	2.421	-	-	-	-	Continuing	Continuing		
Quantity of RDT&E Articles													

A. Mission Description and Budget Item Justification

Funding for this project starts in FY 2013. Original models of Army Black Hawk MEDEVAC helicopters continue to play a major role in maintaining high US troop survival rates in Iraq and Afghanistan by evacuating wounded troops in less than one-hour. In 2009, a VCSA-approved force design update increased the number of air frames in the force from 12 to 15 aircraft for 37 MEDEVAC companies to better meet operational needs. In 2010, the Army Medical Department (AMEDD) accepted life-cycle management of the MEDEVAC MEP from PEO Aviation. In order to achieve required operational capability and enhance commonality across the MEDEVAC fleet, the MEDEVAC MEP program upgrades and retrofits the 256 MEDEVAC legacy helicopters to achieve the medical capability provided by the HH-60M, which is factory built for the MEDEVAC mission.

B. Accomplishments/Planned Programs (\$ in Millions)	FY 2011	FY 2012	FY 2013
Title: MEDEVAC Mission Sensor Forward Looking Infrared Radar (FLIR)	-	-	2.421
Description: MEDEVAC Mission Sensor (MMS) FLIR for UH-60 aircraft. One of the requirements for the UH-60A/L MEDEVAC is a sensor system that will assist the pilots in locating patient pick-up points and assist them in maintaining situational awareness in night and adverse weather conditions. The MMS is currently being qualified for use on the HH-60M aircraft. This system will be installed on UH-60 aircraft using the proven Sponson-Mount FLIR system, which is currently being used in Operation Enduring Freedom (OEF) for the MEDEVAC mission.			
FY 2013 Plans: Will transition from VS7 and complete testing and integration of the Talon FLIR into the aircraft sponson to ensure maximum capability of the sensor, while minimizing impact to aircraft performance.			
Accomplishments/Planned Programs Subtotals	-	-	2.421

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

N/A

D. Acquisition Strategy

Develop in-house or industrial prototypes in government-managed programs to meet military MEDEVAC and regulatory requirements for production and fielding.

Exhibit R-2A, RDT&E Project Justification: PB 2013 Army	DATE: February 2012	
APPROPRIATION/BUDGET ACTIVITY 2040: Research, Development, Test & Evaluation, Army BA 5: Development & Demonstration (SDD)	R-1 ITEM NOMENCLATURE PE 0604807A: Medical Materiel/Medical Biological Defense Equipment - Eng Dev	PROJECT VS8: MEDEVAC MISSION EQUIPMENT PACKAGE (MEP) - END DEV
E. Performance Metrics		
Performance metrics used in the preparation of this justification material	al may be found in the FY 2010 Army Performan	ce Budget Justification Book, dated May 2010.

PE 0604807A: *Medical Materiel/Medical Biological Defense Equipm...* Army

Exhibit R-3, RDT&E Project Cost Analysis: PB 2013 Army

APPROPRIATION/BUDGET ACTIVITY

2040: Research, Development, Test & Evaluation, Army

BA 5: Development & Demonstration (SDD)

R-1 ITEM NOMENCLATURE

PE 0604807A: Medical Materiel/Medical Biological Defense Equipment - Eng Dev

PROJECT

VS8: MEDEVAC MISSION EQUIPMENT

DATE: February 2012

PACKAGE (MEP) - END DEV

Product Development (\$ in Millio	ns)		FY 2	2012	FY 2 Bas			2013 CO	FY 2013 Total			
Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost	Cost To Complete	Total Cost	Target Value of Contract
MEDEVAC Mission Sensor Forward Looking Infrared	TBD	Redstone Arsenal,:AL	-	-		1.800		-		1.800	0.000	1.800	0.000
		Subtotal	-	-		1.800		-		1.800	0.000	1.800	0.000
Support (\$ in Millions)				FY 2	2012	FY 2 Bas			2013 CO	FY 2013 Total			
Cost Category Item	Contract Method & Type	Performing Activity & Location	Total Prior Years Cost	Cost	Award Date	Cost	Award Date	Cost	Award Date	Cost	Cost To Complete	Total Cost	Target Value of Contract
No product/contract costs greater than \$1M individually	SS/UCA	Redstone Arsenal:AL	-	-		0.621		-		0.621	0.000	0.621	0.000
		Subtotal	_	_		0.621		_		0.621	0.000	0.621	0.000

	Total Prior Years			FY 2	013	FY 2	2013	FY 2013	Cost To		Target Value of
	Cost	FY 2	2012	Ва	se	00	co	Total	Complete	Total Cost	Contract
Project Cost Totals	-	-		2.421		-		2.421	0.000	2.421	0.000

Remarks

Exhibit R-4, RDT&E Schedule Profile: PB 2013 Army

APPROPRIATION/BUDGET ACTIVITY

2040: Research, Development, Test & Evaluation, Army
BA 5: Development & Demonstration (SDD)

PACKAGE (MEP) - END DEV

DATE: February 2012

R-1 ITEM NOMENCLATURE
PE 0604807A: Medical Materiel/Medical
Biological Defense Equipment - Eng Dev

PACKAGE (MEP) - END DEV

	FY 2011			FY 2012					FY 2013			FY 2014			FY 2015			FY 2016			FY 2017							
	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
MEDEVAC Mission Sensor (MMS) FLIR																												

Exhibit R-4A, RDT&E Schedule Details: PB 2013 Army	DATE: February 2012	
APPROPRIATION/BUDGET ACTIVITY	R-1 ITEM NOMENCLATURE	PROJECT
2040: Research, Development, Test & Evaluation, Army	PE 0604807A: Medical Materiel/Medical	VS8: MEDEVAC MISSION EQUIPMENT
BA 5: Development & Demonstration (SDD)	Biological Defense Equipment - Eng Dev	PACKAGE (MEP) - END DEV

Schedule Details

	St	art	End				
Events	Quarter	Year	Quarter	Year			
MEDEVAC Mission Sensor (MMS) FLIR	4	2013	4	2013			