

**UNITED STATES
SECURITIES AND EXCHANGE COMMISSION**

Washington, DC 20549

FORM 8-K

CURRENT REPORT

Pursuant to Section 13 or 15(d) of the Securities Exchange Act of 1934

Date of Report (Date of earliest event reported): **August 29, 2008**

PHARMATHENE, INC.

(Exact name of registrant as specified in its charter)

Delaware
(State or other jurisdiction
of incorporation)

001-32587
(Commission
File Number)

20-2726770
(IRS Employer
Identification No.)

One Park Place, Suite 450, Annapolis, Maryland
(Address of principal executive offices)

21401
(Zip Code)

Registrant's telephone number including area code: **(410) 269-2600**

(Former name or former address, if changed since last report)

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:

- Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)
- Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)
- Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))
- Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

Item 7.01. Regulation FD Disclosure

We are furnishing as an exhibit to this report a PowerPoint presentation that representatives of PharmAthene, Inc. (the "Company") plan to use for discussions with certain of the Company's stockholders and other interested persons.

Any information contained in the presentation should be read in the context of and with due regard to the more detailed information provided in other documents we file with or furnish to the Securities and Exchange Commission, including, but not limited to, our annual report on Form 10-K for the year ended December 31, 2007 and our quarterly report on Form 10-Q for the quarter ended June 30, 2008.

Item 9.01. Financial Statements and Exhibits.

(d) Exhibits:

<u>Exhibit No.</u>	<u>Description</u>
99.1	PowerPoint Presentation

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the Registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

PHARMATHENE, INC.
(Registrant)

Date: August 29, 2008

By:

/s/ Christopher C. Camut

Christopher C. Camut

Chief Financial Officer

PharmAthene Investor Presentation

August 29 2008



PharmAthene

Dedicated to a safer world

This presentation contains forward-looking statements within the meaning of Section 27A of the Securities Act of 1933, as amended, and Section 21E of the Securities Exchange Act of 1934, as amended. This information may involve known and unknown risks, uncertainties and other factors that are difficult to predict and may cause the Company's actual results, performance or achievements to be materially different from future results, performance or achievements expressed or implied by any forward-looking statements. Forward-looking statements, which involve assumptions and describe management's current expectations regarding the Company's future plans, strategies and objectives, are generally identifiable by use of the words "may," "will," "should," "expect," "anticipate," "estimate," "believe," "intend," "project," "potential" or "plan" or the negative of these words or other variations on these words or comparable terminology. Such statements include, but are not limited to, statements about future government contract awards, potential payments under government contracts, potential regulatory approvals, future product advancements, anticipated financial results and expected benefits of the acquisition of Avecia Vaccines. These forward-looking statements are based on assumptions that may be incorrect, and we cannot assure you that the projections included in these forward-looking statements will come to pass. The Company's actual results could differ materially from those expressed or implied by the forward-looking statements as a result of various factors, including, but not limited to the "Risk Factors" included in the Company's annual report on Form 10-K and other reports filed with the SEC.

2

2

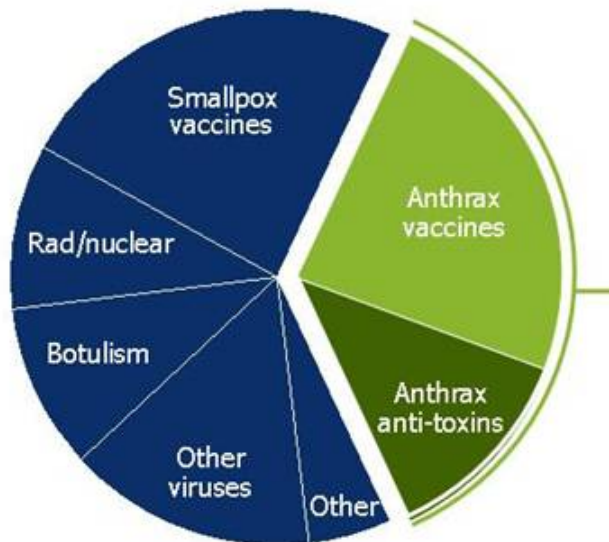
Investment Highlights

- Penetrating multi-billion dollar global biodefense market
- Building highly competitive portfolio of next generation products
- Pursuing focused acquisition strategy with significant revenue opportunities
- Demonstrating strong track record obtaining significant government funding



3

Project BioShield: \$5.6B Market Opportunity

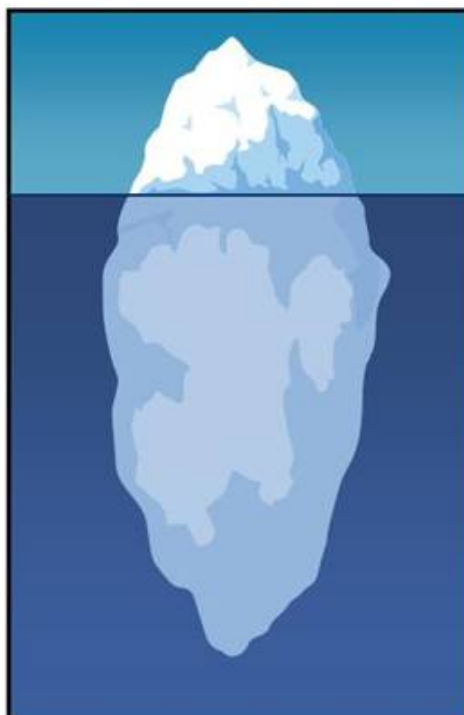


Total anthrax share 36% or \$2B opportunity targeted by PharmAthene



⁴ Source: HHS Public Health Emergency Medical Countermeasure Enterprise Implementation Plan; BioShield contracts awarded

Opportunities Beyond Project BioShield



Project BioShield funding is only the tip of the iceberg

\$5.6B

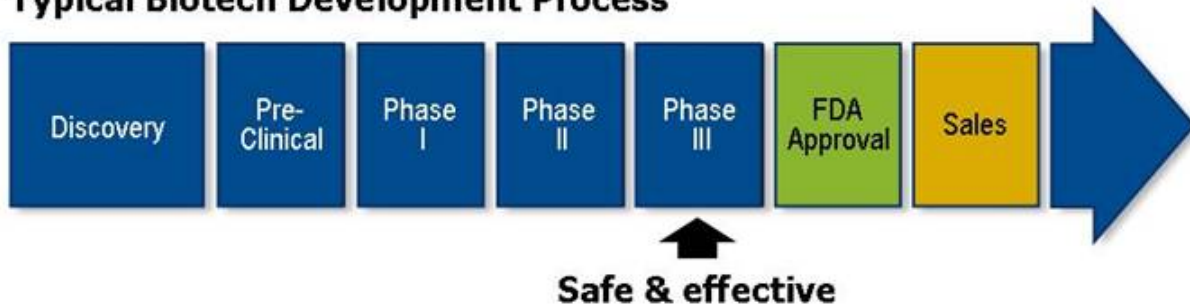
Actively pursuing additional markets:

Department of Defense purchases	\$5B
International purchases	\$6B
Commercial purchases	\$1B
Fortune 500 companies	
Leasing opportunities	
Execution of DHHS Implementation Plan	\$35B

Total biodefense market opportunity ~\$50B

⁵ Source: MedaCorp Reports *Chemical & Biological Defense Program – Oct 2005; DHHS Implementation Plan; Company Estimates through 2018

Typical Biotech Development Process



Biodefense Development Process



- Focus
 - Biodefense market
- Experience
 - Identifying high-priority government needs
 - Identifying and acquiring best-in-class products
 - Collaborating with government to develop and commercialize products
- Success
 - Up to \$485MM* in contracts and funding awarded to date
 - Partner of choice for biodefense



7

**If all milestones are met and options exercised by government*

Executive Leadership

Senior Officers:

David P. Wright
President & Chief Executive Officer

Christopher C. Camut
Chief Financial Officer

Francesca Cook
VP, Policy & Government Affairs

Wayne Morges, PhD
VP, Regulatory Affairs & Quality

Eric I. Richman
SVP, Bus. Dev. & Strategic Planning

Valerie Riddle, MD, FACP
VP and Medical Director

Joan Fusco, PhD
SVP, Operations

Jordan Karp, JD
SVP, General Counsel

Previous Affiliations:

MedImmune, Guilford, GenVec, Smith-Kline & French, G.D. Searle, Glaxo

RecoverCare, Wachovia, Alex Brown & Sons

Guilford, Covance, US Senate, DHHS

Baxter Healthcare, Acambis, Merck

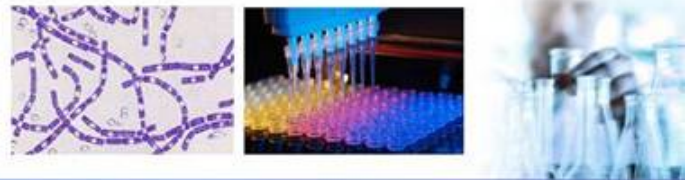
MedImmune, MaxCyte, Healthcare Ventures

MedImmune, Guilford, Washington Hospital

Acambis, Baxter Healthcare

Guilford, Constellation Energy, Mentor, MCI

- Penetrating multi-billion dollar global biodefense market
- Building highly competitive portfolio of next generation products
- Pursuing focused acquisition strategy with significant revenue opportunities
- Demonstrating strong track record obtaining significant government funding



9

Best-in-Class Portfolio

1

rPA Anthrax Vaccine
SparVax™



2

3rd Generation
rPA Anthrax Vaccine



3

Anthrax Anti-Toxin
Valortim®

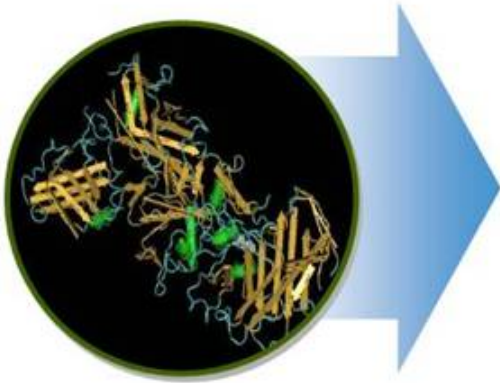


4

Nerve Agent Prophylaxis
Protexia®



Recombinant Protective Antigen (rPA) anthrax vaccine



Characteristics

- Highly purified recombinant version of Protective Antigen
- Produces vaccine-induced antibody response comparable to current licensed vaccine

Advantages

- 3 dose intramuscular regimen vs 6 dose subcutaneous for BioThrax®
- Improved consistency
- Completed Ph II testing in 770 individuals; safe & well tolerated

- rPA Anthrax Vaccine Opportunity
 - Initial DHHS procurement contract (rPA vaccine): 25MM doses
 - USG award date in RFP: December 31, 2008
 - Potential market opportunity in RFP: \$350MM - \$600MM
- Worldwide Anthrax Vaccine Market



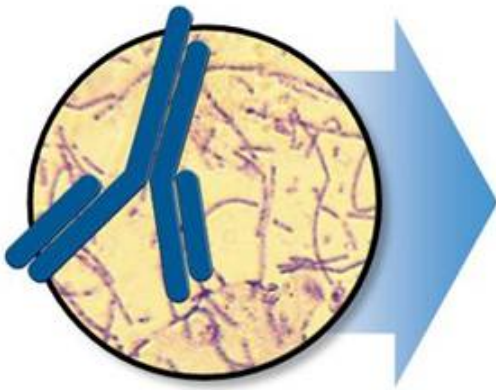
¹² Source: analyst reports; company estimates

3rd Generation rPA Anthrax Vaccine

- Government Requirement
 - Develop 3rd generation rPA-based anthrax vaccine with
 - Enhanced stability – maintain stability for 3 years at 35°C
 - Improved potency – induce protective immunity in 2 or fewer doses
- PharmAthene's 3rd generation product
 - Room temperature stable with enhanced immunogenicity
 - Program funded \$7MM to date by NIH
 - USG award date in RFP, September 2008
- PharmAthene Goal
 - Capture significant market share in both 2nd and 3rd generation vaccine market



Fully human monoclonal antibody (MAb) with a unique mechanism of action



Characteristics

- Fully human monoclonal antibody (MAb)
- Potent anthrax toxin neutralizing activity
- Mechanism of action appears similar to natural immune response

Advantages

- Capable of neutralizing both free and cell-bound anthrax toxin
- Efficacious as both prophylaxis and therapy
- Potential sporicidal activity
- Provides significant, sustained protection to monkeys with a single dose

14

Valortim® - Impressive Data Package

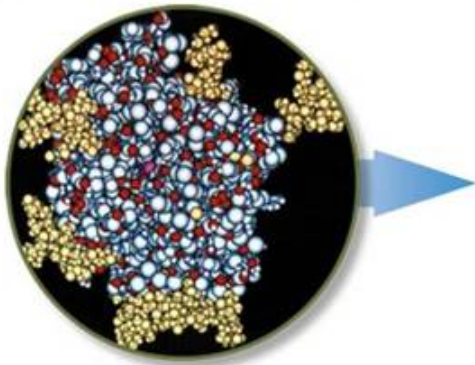
- Initial Phase I in humans complete; no SAE's attributed to Valortim®
- Multiple animal studies have demonstrated efficacy

	Animal	Time to Treatment	Survival
Prophylaxis	Rabbits	1 hr post-exposure	85%
Prophylaxis	Monkeys	1 hr post-exposure	100%
Treatment	Rabbits	24 hrs post-exposure	88%
Treatment	Rabbits	48 hrs post-exposure	42%
Treatment	AG Monkeys	At time of ECL for PA	56%
Control	All Above	All Above	0%

15

- Current options are inadequate
 - Antibiotics are ineffective
 - Vaccines are inappropriate for treatment
- USG requirements established for anti-toxins
 - DHS Material Threat Assessment: 200,000 treatments
 - DHHS procurements to date under 2004 RFP requirement
 - HGSI – 20,000 doses; \$8,260 cost/dose
 - Cangene – 10,000 doses; \$14,383 cost/dose
- Valortim[®] is well positioned for procurement
 - USG funding awarded to date ~\$25MM

Recombinant human BChE (Butyrylcholinesterase)



Mimics natural
"bioscavenger"

Characteristics

- Novel recombinant form of naturally occurring bioscavenger protein
- Produced using innovative transgenic manufacturing platform

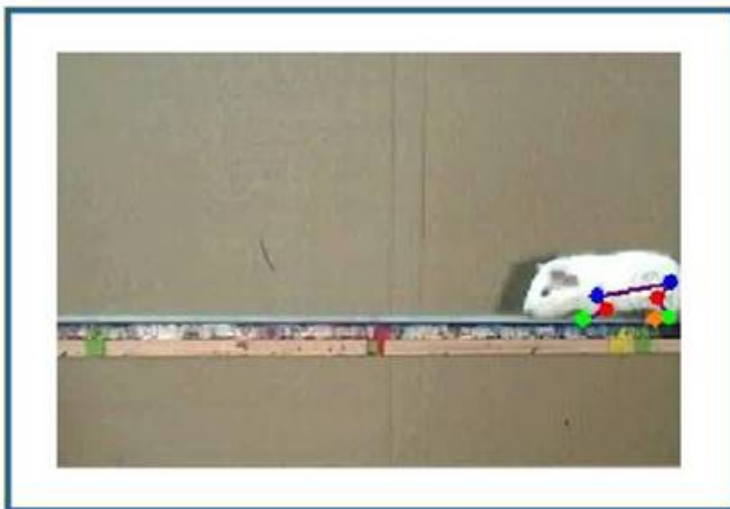
Advantages Over Standard of Care

- Protection pre- and post-exposure
- Protection against broad spectrum of nerve agents
- Superior efficacy to standard of care
- No observable neurological deficits

17

Conventional Treatment Does Not Prevent Neurological Toxicity

Conventional Treatment



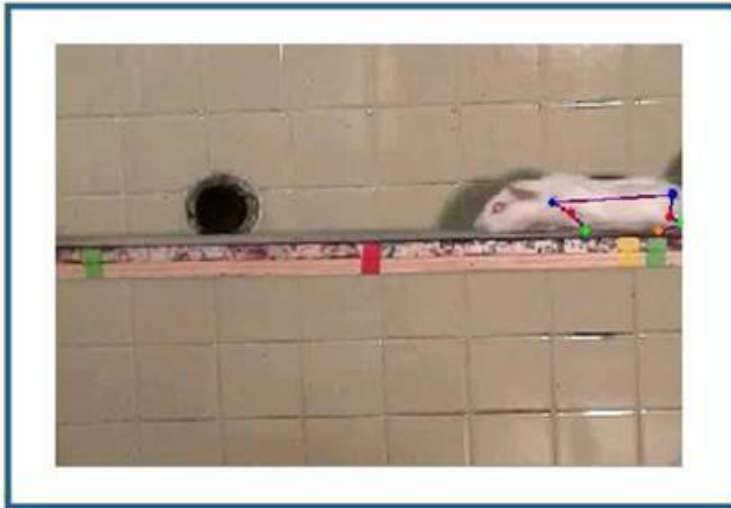
Guinea pig exposed to *only* 1.5 x LD₅₀ Soman and immediately given the conventional treatment of atropine / 2-PAM / Diazepam

- Only 50% of those exposed survived
- Severe neurological deficits

Only Protexia® Provides Superior Survival and Prevents Neurological Toxicity



Protexia® Solution



Guinea pig pretreated with Protexia® and then 18 hours later exposed to 5.5x LD₅₀ of Soman

- 100% survival rate
- No neurological deficits

19 *Source of the film: U.S. Army Medical Research Institute of Chemical Defense*

Nerve Agents - Market Opportunity



- Department of Defense advanced development procurement contract
 - Total value of up to \$219MM*
 - \$106MM in development funding
 - \$113MM for procurement of initial 90,000 doses
- Additional opportunity for civilian (SNS); ex-US military & civilian commercial purchases
- Expanding applications to non-biodefense markets
 - Alzheimer's disease



20 **If all milestones are met and options exercised by the USG.*

- Penetrating multi-billion dollar global biodefense market
- Building highly competitive portfolio of *next generation* products
- Pursuing focused acquisition strategy with significant revenue opportunities
- Demonstrating strong track record obtaining significant government contracts



Unparalleled Track Record in Biodefense



Established strong biodefense portfolio

Building

Significant DoD contract (Protexia®)

up to \$219MM*

Advanced development funding for Valortim®

\$25MM

Includes NIAID/BARDA contract

\$14MM

Total biodefense vaccines government funding

up to \$220MM*

Total amounts under all Government contracts**

up to \$485MM*

22

**If all milestones are met and options exercised by government*

***Includes amounts not set forth above*

Clear Roadmap to Create Value



- 1** Continue to obtain procurement contracts and increase revenues
- 2** Expand portfolio through strategic acquisitions
- 3** Develop multiple government users and non-government customers
- 4** Develop and position products for commercial uses



- Strong cash position
 - \$34.8MM at June 30, 2008
 - > Includes \$15.8MM restricted cash
- Comparatively low cash burn rate
 - Net ~\$1-2MM month, \$4-5MM gross
 - Majority of program development funded with Government grants/contracts
- Substantial ongoing & potential funding contract and grant support
 - Approximately \$60MM development funding for SparVax™ program
 - Opportunity for 3rd generation rPA development contract greater than \$75MM
- Sizeable near-term potential development and procurement contracts*
 - Protexia® procurement contract awarded 2006; up to \$219MM
 - rPA anthrax vaccine procurement contract for 25MM doses: \$350-\$600MM



24 *If all milestones are met and options exercised by the USG.

Current Biodefense Contracts

Product Candidate	Period of Performance	Total Contract Value*	Total Revenue Recognized as of 7/31/08
SparVax™	NIH: 9/03 - 8/13	\$118MM	\$82MM
	NIH: 9/02 - 12/07	\$10.2MM	\$10.2MM
3rd Gen rPA	NIH: 8/05 - 8/10	\$6.9MM	\$3.8MM
Valortim®	NIH: 9/07 - 9/10	\$13.9MM	\$0.6MM
	DoD: 3/07 - 10/09	\$2.7MM	\$0.6MM
	Medarex NIH grants	\$7.2MM	\$7.2MM
Protexia®	DOD: 9/06 - 7/19	\$219MM	\$29.9MM
	NIH: 9/06 - 5/11	\$1.7MM	\$0.1MM
RypVax™	NIH: 9/04 - 9/10	\$50.7MM	\$34.2MM
	DSTL: 5/06 - 6/08	\$40MM	\$37.2MM

25 *Provided certain milestones are achieved and all options are exercised by the government

Key 2008-09 Value Creation Events

	2008		2009	
	H1	H2	H1	H2
Completed strategic acquisition	█			
Completed Valortim® manufacturing scale up	█			
Potential \$75MM 3 rd gen rPA development contract		█		
File Protexia® IND		█		
Begin Protexia® Phase I clinical trial		█		
Potential SparVax™ procurement contract				█
Report Protexia® Phase I results			█	
Potential to begin 2nd phase Protexia® DoD contract (\$64.5MM)				█
Potential advanced funding for anthrax anti-toxin			█	█

- Large and growing market
 - Urgent requirements, multi-billion dollar market
- Experienced management team
 - Previous long-term working relationships with strong execution skills
- Successful execution of growth strategy
 - Advancing three best-in-class, next-generation products
- Solid track record validates our approach
 - Potential government funding/contracts of up to \$485MM* to date
- Clear roadmap for success and value creation



27

*If all milestones are met and options exercised by the USG.

PharmAthene Investor Presentation

August 29 2008



PharmAthene

Dedicated to a safer world