UNITED STATES SECURITIES AND EXCHANGE COMMISSION

Washington, D.C. 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): June 16, 2011

AETHLON MEDICAL, INC.

(Exact name of registrant as specified in its charter)

Nevada	000-21846	13-3632859
(State or other jurisdiction of incorporation)	(Commission File Number)	(I.R.S. Employer Identification No.)
8910 University Center Lane, Suite 660		
San Diego, California		92122
(Address of principal executive offices)		(Zip Code)
Registrant's te	lephone number, including area code: (858) 459-78	800
	Not Applicable	
(Former na	me 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 r	egistrant under any of the following provisions:
Written communications pursuant to Rule 425 under the Securitie	,	
Soliciting material pursuant to Rule 14a-12 under the Exchange A	,	
☐ Pre-commencement communications pursuant to Rule 14d-2(b) u		
Pre-commencement communications pursuant to Rule 13e-4(c) up	nder the Exchange Act (17 CFR 240.13e-4(c))	

Item 7.01 Regulation FD Disclosure

The Registrant disclosed that on Thursday, June 16, 2011, it made a presentation at the LHA Life Sciences & Medical Technology Virtual Conference. A copy of the presentation materials is being furnished as an exhibit to this report and is incorporated by reference into this Item 7.01. Also, the Registrant posted the presentation materials to its website (www.aethlonmedical.com) today, June 16, 2011.

The Registrant notes that the following points made in the presentation were disclosures that may represent new information:

Slide number 5 - The registrant detailed the Aethlon ADAPT(TM) system.

Slide number 11 – The Registrant has initiated a research study to test the ability of the Aethlon Hemopurifier® to capture of HIV related exosomes that transport immunosuppressive NEF protein.

Slide number 29 - The Registrant detailed its government contract and grant submissions.

The Registrant disclosed that it submitted a response to BARDA CBRN BAA-11-100-00009 on May 31, 2011

The Registrant disclosed plans to submit a proposal to U.S. Army, Blood Purification for Organ Failure SBIR by June 29, 2011

Slide number 30 - The registrant disclosed that its wholly owned subsidiary Exosome Sciences, Inc. may license or initiate sales of the ELLSA research diagnostic tools that identify and quantify exosomes. Registrant disclosed that over 70% of researchers polled in market survey indicated interest in purchasing the product.

Slide number 31 - The Registrant provided an update on Hepatitis C clinical studies being conducted at the Medanta-Medicity Institute in Delhi, India.

Slide number 33 – Registrant disclosed clinical program opportunities in the European Union; that it is reviewing an investigator-driven protocol proposal to treat cancer patients on dialysis; and plans to request permission from FDA to initiate U.S. clinical studies.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits. The following exhibit is being furnished pursuant to Item 7.01 above.

Exhibit No.	Description		
99.1	LHA Life Sciences & Medical Technology Virtual Conference Presentation		

SIGNATURES

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

AETHLON MEDICAL, INC.

(Registrant)

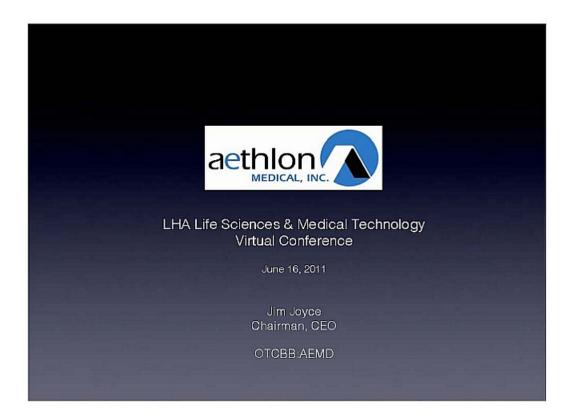
Date: June 16, 2011 By: /s/ James B. Frakes

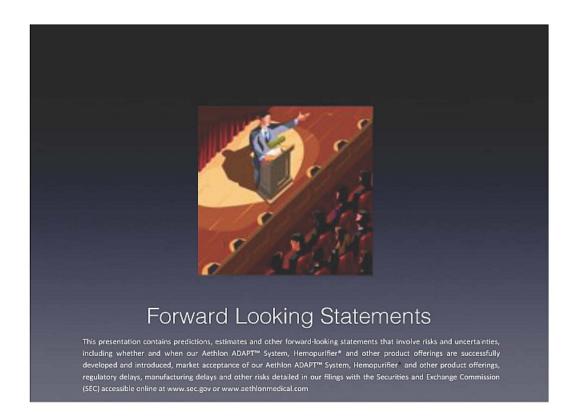
James B. Frakes Chief Financial Officer

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EXHIBIT INDEX

Exhibit No.		Description		
99.1	Presentation Materials			
		4		







Investment Highlights

- · Adaptable Therapeutic Filtration Device Platform
- Our Products Address Unmet Medical Needs
- Our Products Address Large Markets
- Early Adopter Distribution Channel in Place
- · 4 Paths To Early Revenue
- New Clinical Programs



The Aethlon ADAPT™ System

Adaptive Dialysis-like Affinity Platform Technology

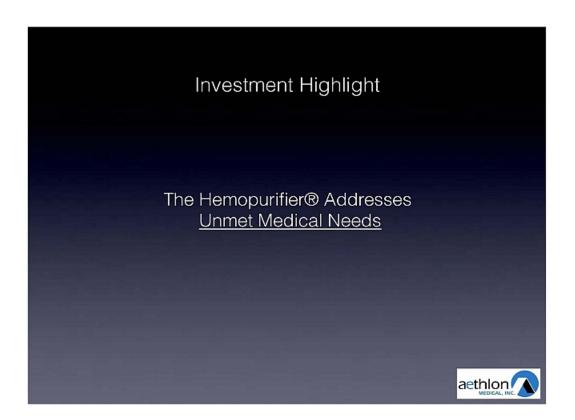
- Converges proven plasma membrane technology with antibodies or other affinity drug agents
- Provides foundation for an expansive pipeline of <u>selective</u> therapeutic filtration (STF) devices
- Transforms drug agent mechanism from biologic to less burdensome medical device regulatory pathway
- Repurposes development-stage, clinical-stage, branded, and clinical dropout affinity drugs

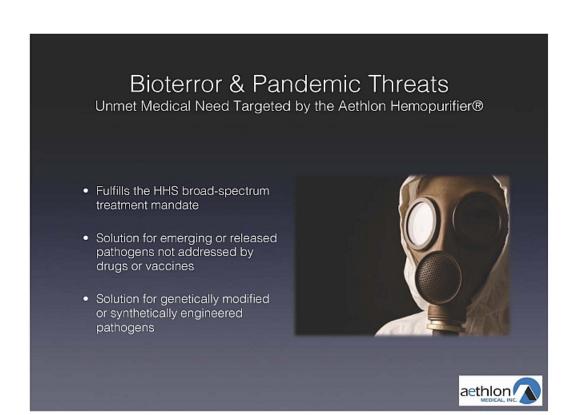


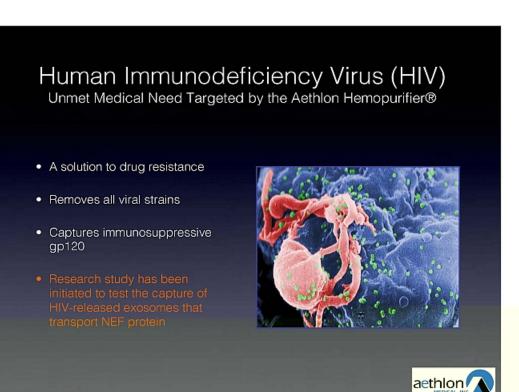
The Adapt™ System Pipeline Introduced to the drug industry on May 25th, 2011 • The Aethlon Hemopurifier® • STF device to reduce incidence of sepsis • Proposed in DARPA Dialysis-Like Therapeutics contract response • STF device to address trauma related organ failure • To be proposed in forthcoming U.S. Army response

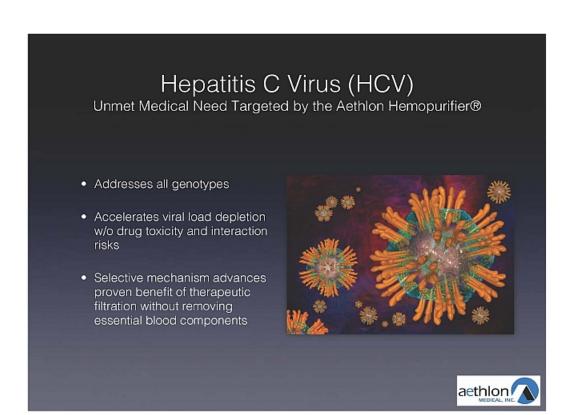


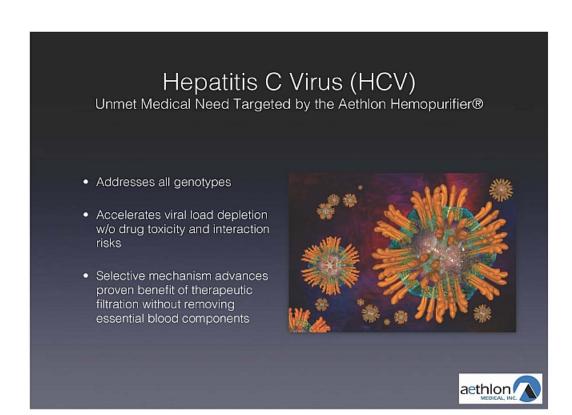








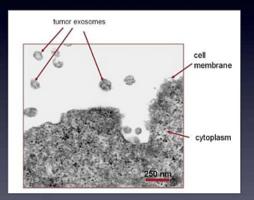




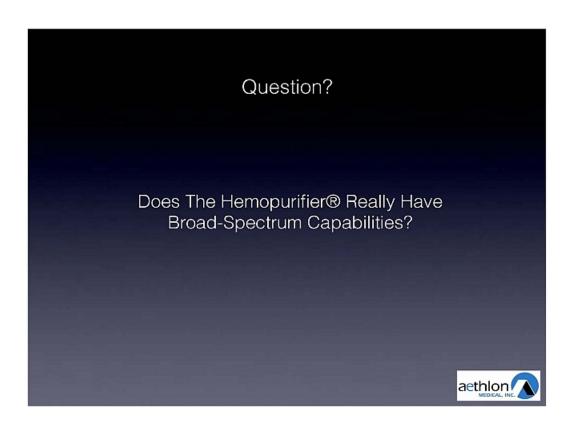
Cancer

Unmet Medical Need Targeted by the Aethlon Hemopurifier®

- The first therapeutic strategy to address exosomes released by cancer
 - Induce apoptosis
 - Disrupt T-cell signaling
 - Inhibit anti-cancer cytokine production
 - Angiogenesis
 - Motaetaeie

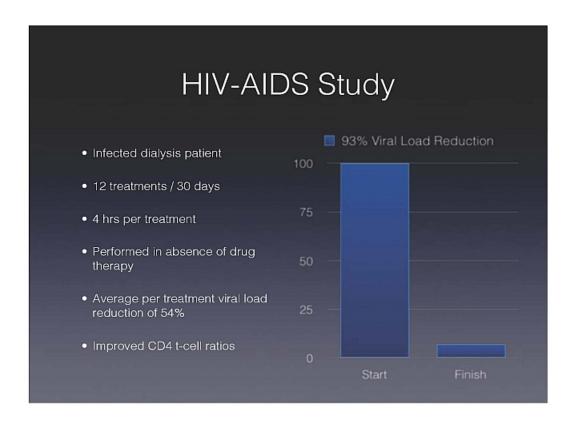


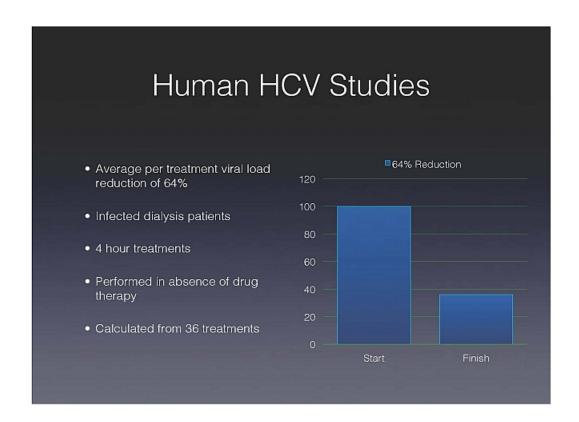




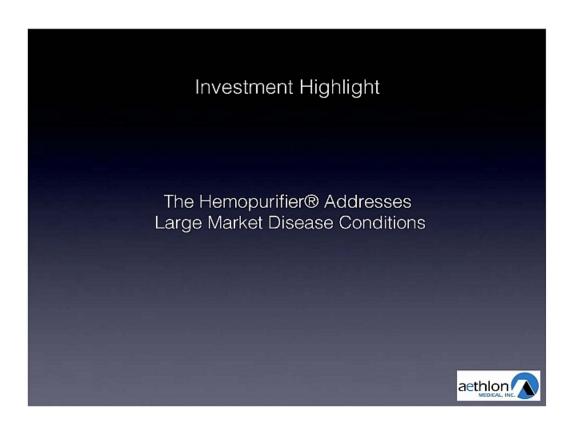
In Vitro Confirmations Against Bioterror and Pandemic Threats

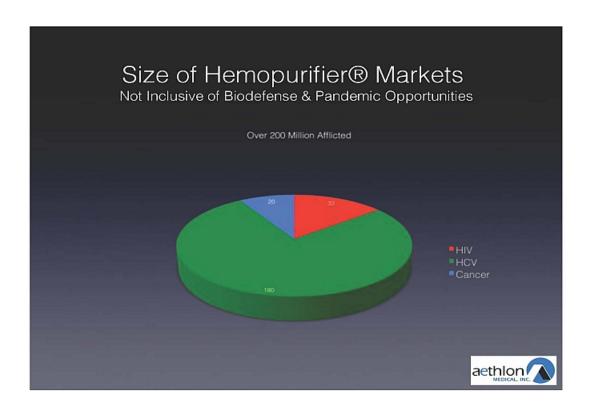
Collaborator	
USAMRIID/CDC	
NIV/WHO	
SFBR	
Battelle	

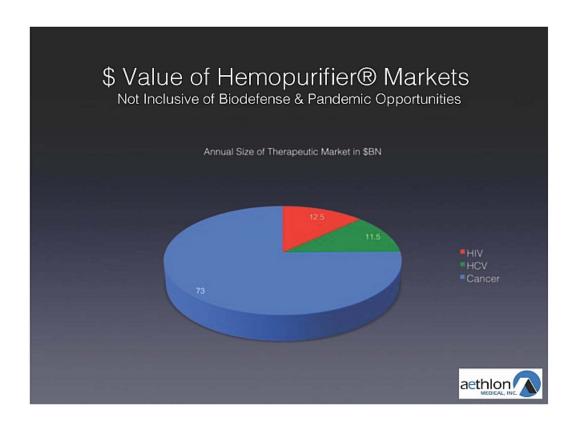




In vitro Cancer Exosome Validations Ovarian Breast Colorectal Lymphoma Melanoma











The Early Adopter Distribution Channel

- 20,000 Nephrologists who will know how to administer Hemopurifier® therapy
- Over 15,000 dialysis centers available to become official Hemopurifier® therapy locations
 - · No additional infrastructure required
- 2 million dialysis patients already being treated by Nephrologists
 - Incidence of HCV is 100-1000x greater than general population
 - · Dialysis patients also have higher incidences of Cancer and HIV
 - Hemopurifier® therapy can be administered during scheduled dialysis treatments







4 Paths to Early Revenue

- 1. Aethlon ADAPT™ system revenues
- 2. Government Contract or Grant Income
- 3. Research Diagnostics
- 4. Hemopurifier® Commericialization



4 Paths to Early Revenue 1.) Aethlon ADAPT™ System Revenues

- Multi-channel revenue opportunity driven by future products created through drug industry collaborations
- Product development fees
- Fees from research, regulatory, and manufacturing support
- Downstream royalties



4 Paths to Early Revenue 2.) Government Contract or Grant Income

- Pending Response to DARPA-BAA-11-30 entitled, "Dialysis-Like Therapeutics" submitted April 1, 2011
- Pending Response to BARDA CBRN BAA-11-100-00009 submitted on May 31, 2011
- Forthcoming submission U.S. Army, Blood Purification for Organ Failure SBIR due June 29th

Reported how displaceurs





4 Paths to Early Revenue 4.) Hemopurifier® Commercialization

- A 30 patient study underway at the Medanta-Medicity Institute in Delhi, India
 - Treatment target: HCV
 - Primary study objective: To demonstrate Hemopurifier® therapy accelerates early viral load depletion when combined with HCV-SOC therapy
 - Achievement of study objectives will trigger first commercialization in India
 - Status 5 patients completed / patient 6 & 7 enrolled
 - Creating a drug subsidy program to accelerate enrollment



Denotes new disclosure



Expansion of Clinical Programs

- Discussing clinical program opportunities in the EU with candidate collaborators and partners
- Reviewing an investigator-driven protocol related to the treatment of cancer patients on dialysis
- We plan to request permission from FDA to initiate U.S. clinical programs this summer
 - IDE on file related to treatment of bioterror and pandemic threats
 - We will also seek permission from FDA to expand label indication to include treatment of HCV

Denotes new disclosure



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