

November 9, 2021



Aethlon Medical Announces Second Quarter Financial Results and Provides Corporate Update

SAN DIEGO, Nov. 9, 2021 /PRNewswire/ -- Aethlon Medical, Inc. (Nasdaq: AEMD), a company developing medical technology to treat cancer and life-threatening infectious diseases, today reported financial results for its second quarter ended September 30, 2021 and provided an update on recent developments.

Company Updates

Aethlon Medical is continuing the research and clinical development of the Hemopurifier™, our therapeutic blood filtration system that can bind and remove life-threatening viruses and harmful exosomes from blood. This action has potential applications in cancer, where cancer associated exosomes may promote immune suppression and metastasis, and in life-threatening infectious diseases, including removal of COVID-19 virus, associated variants, and related exosomes.

As disclosed in our last earnings release on August 9, 2021, the Aethlon Hemopurifier™ has demonstrated binding of SARS-CoV-2 spike protein and, as reported in a peer reviewed publication, the binding and removal from circulation of SARS-CoV-2 virus from a human patient. This is in addition to the Hemopurifier's previously demonstrated binding of numerous pathogenic viruses. The new information about the Hemopurifier in COVID-19 has stimulated clinical researchers to express interest in joining our ongoing clinical trial investigating the Hemopurifier for the treatment of patients with SARS-CoV-2/COVID-19 infection. This trial is being conducted under the open Investigational Device Exemption (IDE) for the Hemopurifier in life threatening viral infections. The trial is designed to allow for up to 40 of these patients to be treated under an Early Feasibility Study protocol at up to 20 clinical sites in the U.S.

During the recent quarter, we entered into an agreement with PPD, Inc., a leading global contract research organization (CRO), to oversee our U.S. clinical studies investigating the Hemopurifier for critically ill COVID-19 patients.

Together with PPD, we continue to advance site readiness at Cooper Medical Center, Loma Linda Medical Center, University of California Davis, Virginia Commonwealth University Medical Center, LSU Health Shreveport, University of Miami Medical Center, and Thomas Jefferson Medical Center. Additionally, we obtained institutional research board approval and have entered into a clinical trial agreement with Stanford Hospital. We are in discussions to bring on board other key U.S. medical centers.

We also recently obtained ethics review board approval and entered into a clinical trial

agreement with Medanta Medicity Hospital, a multi-specialty hospital in Delhi NCR, India, for a COVID-19 clinical trial at that location. On-site training is expected to take place in November 2021.

"The opportunity to help critically ill, ICU patients with COVID-19 continues in both the U.S. and India," said Steven LaRosa, M.D., Chief Medical Officer.

"In addition to our work with COVID-19, we remain very optimistic about the use of our Hemopurifier for the treatment of Head and Neck Cancer. We acknowledge that the enrollment of our Head and Neck Cancer trial has been delayed, primarily due to the COVID-19 pandemic. We are exploring additional avenues to investigate our Hemopurifier in patients with cancer," said Charles J. Fisher, Jr., M.D., CEO.

Financial Results for the Second Quarter Ended September 30, 2021

At September 30, 2021, Aethlon Medical had a cash balance of approximately \$23.2 million.

Aethlon recorded approximately \$115,000 of government contract revenue on its Phase 2 Melanoma Cancer Contract in the three months ended September 30, 2021. We also recorded approximately \$17,000 of revenue related to our cost reimbursable subaward arrangement with the University of Pittsburgh in connection with an NIH contract entitled "Depleting Exosomes to Improve Responses to Immune Therapy in HNNCC." As a result, the Company recorded total government contract revenue of approximately \$132,000 in the three months ended September 30, 2021. Aethlon did not record any government contract revenue in the three months ended September 30, 2020.

Consolidated operating expenses for the three months ended September 30, 2021 were approximately \$2.1 million, compared to approximately \$1.8 million for the three months ended September 30, 2020. This increase of approximately \$300,000, or 20%, in the 2021 period was due to increases in payroll and related expenses of approximately \$200,000 and in general and administrative expenses of approximately \$100,000.

The \$200,000 increase in payroll and related expenses was primarily due to the combination of a \$101,000 increase in our research and development payroll as the result of hiring additional scientists and, a \$100,000 increase in general and administrative payroll expense as the result of additional headcount.

The \$100,000 increase in general and administrative expenses was primarily due to a \$72,000 increase in our rent expense, a \$54,000 increase in our amortization expense and a \$46,000 increase in our insurance expenses, which were partially offset by a \$57,000 decrease in our clinical trial expenses.

As a result of the changes in revenues and expenses noted above, the Company's net loss before noncontrolling interests increased to approximately \$2.0 million for the three months ended September 30, 2021, from approximately \$1.8 million for the three months ended September 30, 2020.

The unaudited condensed consolidated balance sheet for September 30, 2021 and the unaudited condensed consolidated statements of operations for the three and six month periods ended September 30, 2021 and 2020 follow at the end of this release.

Conference Call

The Company will hold a conference call today, Tuesday, November 9, 2021 at 4:30 p.m. Eastern Time to review financial results and recent corporate developments. Following management's formal remarks, there will be a question and answer session.

Interested parties can register for the conference by navigating to <https://dpregrister.com/sreg/10161862/ef905a2ede>.

Please note that registered participants will receive their dial in number upon registration.

Interested parties without internet access or unable to pre-register may dial in by calling:
PARTICIPANT DIAL IN (TOLL FREE): 1-844-836-8741
PARTICIPANT INTERNATIONAL DIAL IN: 1-412-317-5442

All callers should ask for the Aethlon Medical, Inc. conference call.

A replay of the call will be available approximately one hour after the end of the call through December 9, 2021. The replay can be accessed via Aethlon Medical's website or by dialing 1-877-344-7529 (domestic) or 1-412-317-0088 (international) or Canada Toll Free at 1-855-669-9658. The replay conference ID number is 10162862.

About Aethlon and the Hemopurifier®

Aethlon Medical is a biotechnology company developing the Hemopurifier, a therapeutic blood filtration system indicated for infectious diseases and cancer. In human studies, the Hemopurifier has demonstrated the removal of life-threatening viruses and harmful exosomes from blood utilizing a proprietary lectin-based technology. This action has potential applications in cancer, where exosomes may promote immune suppression and metastasis, and in life-threatening infectious diseases.

The Hemopurifier is a U.S. Food and Drug Administration (FDA) designated Breakthrough Device indicated for the treatment of individuals with advanced or metastatic cancer who are either unresponsive to or intolerant of standard of care therapy, and with cancer types in which exosomes have been shown to participate in the development or severity of the disease. Under an Investigational Device Exemption (IDE) application, the FDA approved a single site, open-label Early Feasibility Study (EFS) to evaluate the Hemopurifier for reducing cancer-associated exosomes prior to the administration of standard-of-care pembrolizumab (KEYTRUDA®) in patients with recurrent and/or metastatic squamous cell carcinoma of the head and neck. The EFS is being conducted at the University of Pittsburgh Medical Center Hillman Cancer Center.

The Hemopurifier also holds an FDA Breakthrough Device designation and an open IDE application related to the treatment of life-threatening viruses that are not addressed with approved therapies. A recent amendment to the IDE enabled Aethlon to implement a new EFS protocol to treat up to 40 COVID-19 patients at up to 20 clinical sites in the U.S. In two case studies of patients treated under Emergency Use (EU), the Hemopurifier demonstrated binding of SARS-CoV-2 spike protein and removal of SARS-CoV-2 virus from the circulation of a human patient.

Additional information can be found at www.AethlonMedical.com.

This press release contains forward-looking statements within the meaning of Section 27A of the Securities Act of 1933 and Section 21E of the Securities Exchange Act of 1934 that involve risks and uncertainties. Statements containing words such as "may," "believe," "anticipate," "expect," "intend," "plan," "project," "will," "projections," "estimate," "potentially" or similar expressions constitute forward-looking statements. Such forward-looking statements are subject to significant risks and uncertainties and actual results may differ materially from the results anticipated in the forward-looking statements. These forward-looking statements are based upon Aethlon's current expectations and involve assumptions that may never materialize or may prove to be incorrect. Factors that may contribute to such differences include, without limitation, the Company's ability to enroll additional sites for its clinical trials, the Company's ability to enroll patients in and successfully complete its trials in COVID-19 patients and in its head and neck cancer trials, the Company's ability to successfully treat patients under any Emergency Use pathway, the Company's ability to successfully complete development of its Hemopurifier, and other potential risks. The foregoing list of risks and uncertainties is illustrative but is not exhaustive. Additional factors that could cause results to differ materially from those anticipated in forward-looking statements can be found under the caption "Risk Factors" in the Company's Annual Report on Form 10-K for the year ended March 31, 2021, and in the Company's other filings with the Securities and Exchange Commission, including its quarterly Reports on Form 10-Q. All forward-looking statements contained in this press release speak only as of the date on which they were made. Except as may be required by law, the Company does not intend, nor does it undertake any duty, to update this information to reflect future events or circumstances.

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AETHLON MEDICAL, INC. AND SUBSIDIARIES
Condensed Consolidated Balance Sheet

ASSETS	September 30, 2021	March 31, 2021
CURRENT ASSETS		
Cash	\$23,224,925	\$9,861,575
Accounts receivable	131,966	149,082
Prepaid expenses	212,308	341,081
	23,569,199	10,351,738
TOTAL CURRENT ASSETS		
Property and equipment, net	213,625	160,976
Right-of-use lease asset	-	40,363
Patents, net	2,476	56,954
Restricted cash	46,726	46,726
Deposits	42,159	12,159
	238,975	217,278
TOTAL ASSETS		
	\$23,874,185	\$10,668,916
LIABILITIES AND STOCKHOLDERS' EQUITY		
CURRENT LIABILITIES		
Accounts payable	254,176	337,678
Due to related parties	134,207	118,520
Deferred revenue	114,849	114,849
Lease liability	-	42,543
Other current liabilities	471,117	761,636
	974,349	1,375,226
TOTAL CURRENT LIABILITIES		
	974,349	1,375,226
TOTAL LIABILITIES		
COMMITMENTS AND CONTINGENCIES		
EQUITY		
Common stock, par value of \$0.001, 30,000,000 shares authorized; 15,397,299 and 12,150,597 issued and outstanding	15,399	12,152
Additional-paid in capital	147,041,683	129,331,542
Accumulated deficit	(124,018,372)	(119,913,090)
	23,038,710	9,430,604
TOTAL STOCKHOLDERS' EQUITY BEFORE NONCONTROLLING INTERESTS		
Noncontrolling interests	(138,874)	(136,914)
	22,899,836	9,293,690
TOTAL STOCKHOLDERS' EQUITY		
	22,899,836	9,293,690
TOTAL LIABILITIES AND STOCKHOLDERS' EQUITY		
	\$23,874,185	\$10,668,916

AETHLON MEDICAL, INC. AND SUBSIDIARIES
Condensed Consolidated Statements of Operations
For the three and six months ended September 30, 2021 and 2020

	Three Months Ended 9/30/21	Three Months Ended 9/30/20	Six Months Ended 9/30/21	Six Months Ended 9/30/20
Government contract revenue	\$131,966	\$-	\$263,932	\$-
OPERATING COSTS AND EXPENSES				
Professional fees	649,460	656,396	1,232,929	1,220,680
Payroll and related	805,608	560,244	1,822,350	997,155
General and administrative	685,702	554,749	1,315,895	964,700
	<u>2,140,770</u>	<u>1,771,389</u>	<u>4,371,174</u>	<u>3,182,535</u>
OPERATING LOSS	(2,008,804)	(1,771,389)	(4,107,242)	(3,182,535)
NET LOSS	\$(2,008,804)	\$(1,771,389)	\$(4,107,242)	\$(3,182,535)
Loss attributable to noncontrolling interests	(825)	(825)	(1,960)	(1,688)
NET LOSS ATTRIBUTABLE TO AETHLON MEDICAL, INC.	<u>\$(2,007,979)</u>	<u>\$(1,770,564)</u>	<u>\$(4,105,282)</u>	<u>\$(3,180,847)</u>
Basic and diluted net loss available to common stockholders per share	\$ (0.13)	\$ (0.15)	\$ (0.29)	\$ (0.29)
Weighted average number of common shares outstanding	<u>15,386,486</u>	<u>12,070,592</u>	<u>14,114,639</u>	<u>10,845,049</u>

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