
**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: January 29, 2009
(Date of earliest event reported)

ENTEROMEDICS INC.

(Exact name of registrant as specified in its charter)

Commission File Number: 1-33818

Delaware
(State or other jurisdiction of incorporation)

48-1293684
(IRS Employer Identification No.)

2800 Patton Road, St. Paul, Minnesota 55113
(Address of principal executive offices, including zip code)

(651) 634-3003
(Registrant's telephone number, including area code)

Not Applicable
(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))
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Item 2.02. Results of Operations and Financial Condition.

Furnished herewith as Exhibit 99.1 and incorporated by reference herein is the text of EnteroMedics Inc.'s press release regarding its financial results for the year ended December 31, 2008.

The information furnished herewith pursuant to Item 2.02 of this Current Report and in Exhibit 99.1 hereto is being "furnished" in accordance with General Instruction B.2 of Form 8-K and shall not be deemed "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the "Exchange Act"), or otherwise subject to the liabilities of that section, nor shall it be deemed to be incorporated by reference in any filing under the Securities Act of 1933, as amended, or the Exchange Act, whether made before or after the date hereof, regardless of any general incorporation language in such filing.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits

<u>Exhibit No.</u>	<u>Description</u>
99.1	Press release, dated January 29, 2009, entitled "EnteroMedics Reports 2008 Year-End Financial Results."

SIGNATURE

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.

ENTEROMEDICS INC.

By: /s/ Greg S. Lea
Greg S. Lea
Senior Vice President and Chief
Financial Officer

Date: January 29, 2009

EXHIBIT INDEX

<u>Exhibit No.</u>	<u>Description</u>
99.1	Press release, dated January 29, 2009, entitled "EnteroMedics Reports 2008 Year-End Financial Results."



Contact:
EnteroMedics Inc.
Greg S. Lea
(651) 789-2860
ir@enteromedics.com

EnteroMedics Reports 2008 Year-End Financial Results

ST. PAUL, Minn., January 29, 2009 — EnteroMedics Inc., (NASDAQ: ETRM), the developer of medical devices using neuroblocking technology to treat obesity, its associated co-morbidities, and other gastrointestinal disorders, today announced financial results for the three months and full year ended December 31, 2008.

For the full year 2008, the Company reported a net loss of \$37.9 million, or \$2.25 per share, research and development expenses of \$27.7 million, and general and administrative expenses of \$8.6 million. For the fourth quarter ended December 31, 2008, the Company reported a net loss of \$7.8 million, or \$0.46 per share. Expenses were primarily associated with the cost of supporting the Company's multiple ongoing clinical trials as well as the continued development of VBLOC™ vagal blocking therapy (VBLOC Therapy) delivered through the Company's Maestro™ System. On December 31, 2008, the Company's cash, cash equivalents and short-term investments totaled \$26.3 million.

"In 2008, the Company continued to acquire data that support the use of its technology in obesity as well as the early, yet significant, effects of VBLOC Therapy on the co-morbidities of diabetes and hypertension," said President and CEO Mark B. Knudson, Ph.D. "These results speak to the potential of this technology to directly address the healthcare costs associated with obesity while helping people achieve significant weight loss. We believe that 2009 will see the company achieve a number of key milestones, including further data in Type 2 diabetes and hypertension, and the unblinding of our pivotal EMPOWER trial, the results of which, if favorable, will support our 2009 PMA submission to the FDA for marketing approval of the Maestro System."

Gregory S. Lea, Senior Vice President and Chief Financial Officer of EnteroMedics added: "With the EMPOWER trial fully enrolled and all subjects implanted, a significant portion of the expense associated with that study is now behind us." He concluded, "In addition, we are projecting that our current cash position is sufficient to fund our comprehensive development and regulatory efforts for 2009."

About EnteroMedics Inc.

EnteroMedics is a development stage medical device company focused on the design and development of devices that use neuroblocking technology to treat obesity and other gastrointestinal disorders. EnteroMedics' proprietary neuroblocking technology, VBLOC™ vagal blocking therapy, is designed to

intermittently block the vagus nerves using high-frequency, low-energy, electrical impulses. EnteroMedics has met its enrollment goal under an FDA-approved Investigational Device Exemption (IDE) for the EMPOWER Study using the Maestro™ System, its initial product for the treatment of obesity. EnteroMedics is currently recruiting patients outside of the United States for a feasibility study examining VBLOC Therapy's effects on blood glucose levels in diabetic patients. For more information, visit www.enteromedics.com.

Forward-Looking Safe Harbor Statement:

This press release contains forward-looking statements about EnteroMedics Inc. Our actual results could differ materially from those discussed due to known and unknown risks, uncertainties and other factors including our limited history of operations, our losses since inception and for the foreseeable future; our lack of regulatory approval for our Maestro™ System for the treatment of obesity; our inability to complete our EMPOWER pivotal trial and other clinical trials, or significant delays in the completion of our clinical trials; our ability to timely commercialize our Maestro System; our dependence on third parties to initiate and perform our clinical trials; the need to obtain regulatory approval for any modifications to our Maestro System; physician adoption of our Maestro System and VBLOC™ vagal blocking therapy; our ability to obtain third party coding, coverage or payment levels; ongoing regulatory compliance; our dependence on third party manufacturers and suppliers; the successful development of our sales and marketing capabilities; our ability to raise additional capital when needed; our ability to attract and retain management and other personnel and to manage our growth effectively; potential product liability claims; potential healthcare fraud and abuse claims; and our ability to obtain and maintain intellectual property protection for our technology and products. These and additional risks and uncertainties are described more fully in the Company's filings with the Securities and Exchange Commission, particularly those factors identified as "risk factors" in the Company's Form 10-K dated March 13, 2008. We are providing this information as of the date of this press release and do not undertake any obligation to update any forward-looking statements contained in this document as a result of new information, future events or otherwise.

Caution-Investigational device. Limited by Federal law to investigational use.

The implantation procedure and usage of the Maestro™ System carry some risks, such as the risk generally associated with laparoscopic procedures and those related to treatment as described in the EMPOWER clinical trial informed consent.

(See Attached Table)

ENTEROMEDICS INC.
(A Development Stage Company)

Condensed Consolidated Statements of Operations (unaudited)
(in thousands, except per share data)

	Three Months Ended		Year Ended	
	December 31,		December 31,	
	2008	2007	2008	2007
Operating expenses:				
Research and development	\$ 4,387	\$ 7,159	\$ 27,674	\$ 21,053
Selling, general and administrative	2,081	1,510	8,597	6,973
Total operating expenses	6,468	8,669	36,270	28,026
Loss from operations	(6,468)	(8,669)	(36,270)	(28,026)
Other expense, net	(1,354)	(77)	(1,604)	(549)
Net loss	<u>\$ (7,822)</u>	<u>\$ (8,746)</u>	<u>\$ (37,874)</u>	<u>\$ (28,575)</u>
Net loss per share - basic and diluted	<u>\$ (0.46)</u>	<u>\$ (1.10)</u>	<u>\$ (2.25)</u>	<u>\$ (11.69)</u>
Shares used to compute basic and diluted net loss per share	<u>16,879</u>	<u>7,918</u>	<u>16,836</u>	<u>2,445</u>

ENTEROMEDICS INC.
(A Development Stage Company)

Condensed Consolidated Balance Sheets (unaudited)
(in thousands)

	December 31, 2008	December 31, 2007
ASSETS		
Cash, cash equivalents and short-term investments	\$ 26,295	\$ 57,031
Prepaid expenses and other current assets	499	523
Property and equipment, net	1,264	1,492
Other assets	221	5
Total assets	<u>\$ 28,279</u>	<u>\$ 59,051</u>
LIABILITIES AND STOCKHOLDERS' EQUITY		
Liabilities:		
Accounts payable	\$ 163	\$ 300
Debt	13,670	11,099
Other liabilities	3,040	2,370
Total liabilities	<u>16,874</u>	<u>13,769</u>
Stockholders' equity	<u>11,405</u>	<u>45,282</u>
Total liabilities and stockholders' equity	<u>\$ 28,279</u>	<u>\$ 59,051</u>

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