

**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): September 15, 2022

MICROBOT MEDICAL INC.
(Exact name of registrant as specified in its charter)

Delaware
(State or other jurisdiction
of incorporation)

000-19871
(Commission
File Number)

94-3078125
(IRS Employer
Identification No.)

25 Recreation Park Drive, Unit 108
Hingham, Massachusetts 02043
(Address of Principal Executive Offices) (Zip Code)

Registrant's telephone number, including area code: (781) 875-3605

(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))

Securities registered pursuant to Section 12(b) of the Act:

| <u>Title of each class</u> | <u>Trading Symbol(s)</u> | <u>Name of each exchange on which registered</u> |
|--------------------------------|--------------------------|--|
| Common Stock, \$0.01 par value | MBOT | NASDAQ Capital Market |

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (17 CFR §230.405) or Rule 12b-2 of the Securities Exchange Act of 1934 (17 CFR §240.12b-2).

Emerging Growth Company

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act.

Item 7.01 Regulation FD Disclosure.

On September 15, 2022, Microbot Medical Inc. (the “Company”) issued a press release announcing that Vincent Vidal, M.D., PhD, has joined its Scientific Advisory Board (SAB).

The press release, which is furnished as Exhibit 99.1 to this Current Report on Form 8-K, is incorporated herein by reference. The information in this report (including Exhibit 99.1) is being furnished pursuant to Item 7.01 and shall not be deemed to be “filed” for the purposes of Section 18 of the Securities Exchange Act of 1934, as amended, or otherwise subject to the liabilities of that section. This report will not be deemed an admission as to the materiality of any information herein (including Exhibit 99.1).

Item 9.01. Financial Statements and Exhibits.*(d) Exhibits*

| Exhibit Number | Description |
|---------------------------|---|
| 99.1 | Press Release |
| 104 | Cover Page Interactive Data File (embedded within the Inline XBRL document) |

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 thereunto duly authorized.

MICROBOT MEDICAL INC.

By: /s/ Harel Gadot

Name: Harel Gadot

Title: Chief Executive Officer, President and Chairman

Date: September 15, 2022



Microbot Medical Enhances its Scientific Advisory Board (SAB) with its Newest Member Dr. Vincent Vidal

Prominent France based interventional radiologist to leverage his expertise to assist the Company to achieve its future objective for the LIBERTY[®] Robotic System

HINGHAM, Mass., September 15, 2022 – Microbot Medical Inc. (Nasdaq: MBOT), which continues to build a network of global leaders in the endovascular interventional space for the LIBERTY[®] Robotic System, today announced that Vincent Vidal, M.D., PhD, has joined its Scientific Advisory Board (SAB).

Dr. Vidal is a world-leading advocate and distinguished expert of embolization techniques. He is the head of Interventional Radiology in the Imaging Department of Timone Hospital (Marseille, France) and Professor at the Faculty of Medicine and Director at the Experimental Interventional Imaging Laboratory (LiiE) at the Aix-Marseille University, where his research and clinical work are focused on embolization. Dr. Vidal received both his M.D. and PhD from the School of Medicine of Aix-Marseille University, and he is an active member of SIR, CIRSE, ESR, and APSCVIR.

Recognizing the future benefits that the LIBERTY Robotic System can bring to his patients, Dr. Vidal led a successful feasibility pre-clinical trial for the LIBERTY Robotic System at his lab located at LiiE, and the positive results of the trial prompted him to join the Company's SAB where his experience and expertise can be leveraged to help ensure the LIBERTY Robotic System is available to meet the future needs of the interventional radiologist community and their patients.

"We welcome Dr. Vidal as a global leader in the interventional radiology community, and his addition to our SAB is yet another milestone in our vision to allow access to the benefits that the LIBERTY System is expected to deliver to many patients and their healthcare providers globally," commented Harel Gadot, Chairman, CEO, and President of Microbot Medical. "It is always reassuring when top Key Opinion Leaders like Dr. Vidal are excited to join our SAB after their personal experience using our technology, and we believe that the team of scientific leaders we established over the last few months will help position us for a successful regulatory process, both for FDA and CE, and eventually to lead to global commercialization of the world's first fully disposable endovascular robotic system."

"I was immediately drawn to the LIBERTY Robotic System due to its novel form factor and unique capabilities, and once I was able to experience it first-hand, it confirmed my initial views of its future potential," Commented Dr. Vidal. "The Microbot Medical team has achieved a great deal getting it to this point, and my goal is to ensure success at each of the next regulatory steps. I believe my fellow interventional radiologists, as well as the hospital administrators, will gravitate to the LIBERTY Robotic System when it becomes commercially available because I believe it addresses many clinical and financial needs."

About Microbot Medical

Microbot Medical Inc. (NASDAQ: MBOT) is a pre-clinical medical device company that specializes in transformational micro-robotic technologies, focused primarily on both natural and artificial lumens within the human body. Microbot's current proprietary technological platforms provide the foundation for the development of a Multi Generation Pipeline Portfolio (MGPP).

Microbot Medical was founded in 2010 by Harel Gadot, Prof. Moshe Shoham, and Yossi Bornstein with the goals of improving clinical outcomes for patients and increasing accessibility through the use of micro-robotic technologies. Further information about Microbot Medical is available at <http://www.microbotmedical.com>.

Safe Harbor

Statements to future financial and/or operating results, future growth in research, technology, clinical development, and potential opportunities for Microbot Medical Inc. and its subsidiaries, along with other statements about the future expectations, beliefs, goals, plans, or prospects expressed by management, constitute forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995 and the Federal securities laws. Any statements that are not historical fact (including, but not limited to statements that contain words such as "will," "believes," "plans," "anticipates," "expects" and "estimates") should also be considered to be forward-looking statements. Forward-looking statements involve risks and uncertainties, including, without limitation, market conditions, risks inherent in the development and/or commercialization of potential products, including LIBERTY and SCS, the outcome of its studies to evaluate LIBERTY, SCS and other existing and future technologies, any failure or inability to recruit physicians and clinicians to serve as primary investigators to conduct the SCS's early feasibility study which could adversely affect or delay such study, uncertainty in the results of pre-clinical and clinical trials or regulatory pathways and regulatory approvals, uncertainty resulting from the COVID-19 pandemic, need and ability to obtain future capital, and maintenance of intellectual property rights. Additional information on risks facing Microbot Medical can be found under the heading "Risk Factors" in Microbot Medical's periodic reports filed with the Securities and Exchange Commission (SEC), which are available on the SEC's web site at www.sec.gov. Microbot Medical disclaims any intent or obligation to update these forward-looking statements, except as required by law.

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