

**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 9, 2025

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.)

175 Derby St., Bld. 27
Hingham, MA 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:

Title of each class	Trading Symbol(s)	Name of each exchange on which registered
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 June 9, 2025, Microbot Medical Inc. (the “Company”) issued a press release announcing the continued expansion of its commercial team with the addition of Michael Lytle as the head of Sales Operations & Analytics, in preparation for the anticipated U.S. launch of the LIBERTY System, which is projected during the third quarter of 2025. The press release further announced that the Company remains engaged with the U.S. Food and Drug Administration (“FDA”), with a 510(k) decision with respect to its LIBERTY[®] Endovascular Robotic System now expected during the third quarter of 2025. This updated FDA decision timeline remains within the FDA’s original scheduled review window and is not expected to affect the Company’s planned launch upon clearance.

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 Item 7.01 and Exhibit 99.1 is being furnished 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 in this Item 7.01 or Exhibit 99.1.

Item 8.01 Other Events.

The Company announced that a 510(k) decision with respect to its LIBERTY[®] Endovascular Robotic System is now expected during the third quarter of 2025. This updated FDA decision timeline remains within the FDA’s original scheduled review window and is not expected to affect the Company’s planned launch upon clearance

Forward Looking Statements

This Item 8.01 of this Current Report on Form 8-K may contain “forward-looking statements.” Such statements which are not purely historical (including, but not limited to statements that contain words such as “will,” “believes,” “plans,” “anticipates,” “expects,” “intends,” “would,” “could” and “estimates”) are forward-looking statements and include any statements regarding beliefs, plans, expectations or intentions regarding the future, including but not limited to, regulatory milestones.

Actual results could differ from those projected in any forward-looking statements due to numerous factors. These forward-looking statements are made as of the date of this Form 8-K, and the Company assumes no obligation to update the forward-looking statements, or to update the reasons why actual results could differ from those projected in the forward-looking statements, except as required by law. Although the Company believes that the beliefs, plans, expectations and intentions contained in this Form 8-K are reasonable, there can be no assurance that such beliefs, plans, expectations or intentions will prove to be accurate. Investors should consult all of the information set forth herein and should also refer to the risk factors disclosure outlined in the Company’s reports and statements filed from time-to-time with the Securities and Exchange Commission.

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

Exhibit Number	Description
99.1	Press release dated June 9, 2025
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: June 9, 2025

Microbot Medical Continues to Strengthen Commercial Capabilities in Preparation for the anticipated Q3 2025 Launch of its LIBERTY® Endovascular Robotic System

The addition of the head of Sales Operations & Analytics aims to strengthen sales infrastructure and launch execution

HINGHAM, Mass., June 9, 2025 – Microbot Medical Inc. (Nasdaq: MBOT), developer of the innovative LIBERTY Endovascular Robotic System, today announced the continued expansion of its commercial team in preparation for the anticipated U.S. launch of the LIBERTY System, which is projected during the third quarter of 2025. The Company remains actively engaged with the U.S. Food and Drug Administration (FDA), with a 510(k) decision now expected during the third quarter of this year. This updated FDA decision timeline remains within the FDA’s original scheduled review window and is not expected to affect the Company’s planned launch upon clearance.

“Our recent interactions with the FDA have been productive, and we remain confident in the process and outcome,” said Harel Gadot, Chairman, CEO & President. “We continue to operate with full momentum and position the Company for an anticipated launch during the third quarter of this year, which includes the continued build out of our commercial infrastructure and preparing the organization to deliver on our strategy.”

As part of the Company’s ongoing launch preparation, it recently expanded its commercial team with the addition of **Michael Lytle** as the head of Sales Operations & Analytics. Mr. Lytle brings deep experience in sales support, data analysis, and operational excellence. He is expected to play a key role in shaping market intelligence tools and processes to optimize the sales cycle and drive strategic growth.

Prior to joining Microbot, Mr. Lytle held increasing leadership roles at ZOLL Cardiac Management Solutions (CMS), a division of ZOLL Medical Corporation, where he supported the rollout of innovative cardiac care technologies.

“We believe that Michael’s addition will strengthen our launch readiness plans, and that his expertise will help us target the right markets, allocate resources effectively, and accelerate commercial execution,” concluded Mr. Gadot.

The LIBERTY System is currently under FDA review, is not available for sales in the US, and clearance is not guaranteed.

About Microbot Medical

Microbot Medical Inc. (NASDAQ: MBOT) is a pre-commercial stage medical technology company with a vision to redefine endovascular robotics and improve the quality of care for millions of patients and providers globally. The Company has developed the world’s first single-use, fully disposable endovascular robotic system, which aims to eliminate traditional barriers to accessing advanced robotic systems.

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, commercialization 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, the Company’s need for and ability to obtain additional working capital to continue its transition to a commercially focused company, market conditions, risks inherent in the development and/or commercialization of the LIBERTY® Endovascular Robotic System, uncertainty in the results of regulatory pathways and regulatory approvals, including whether the FDA will timely grant 510(k) clearance to commercially market the LIBERTY® Endovascular Robotic System in the United States if at all, uncertainty resulting from political, social and geopolitical conditions, particularly any changes in personnel or processes or procedures at the FDA and announcements of tariffs on imports into the U.S., disruptions resulting from new and ongoing hostilities between Israel and the Palestinians and other neighboring countries, 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

Investor Contact:

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