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): October 22, 2024

CYCLACEL PHARMACEUTICALS, INC.

(Exact name of registrant as specified in its charter)

Delaware (State or other jurisdiction of incorporation) 0-50626 (Commission File Number) 91-1707622 (IRS Employer Identification No.)

200 Connell Drive, Suite 1500 Berkeley Heights, NJ 07922 (Address of principal executive offices and zip code)

Registrant's telephone number, including area code: (908) 517-7330

(Former Name or Former Address, if Changed Since Last Report)

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Check the appropriate box below if the Form 8-K f following provisions (<i>see</i> General Instruction A.2. b	e	sly satisfy the filing obligation of the registrant under any of the
☐ Written communications pursuant to Rule 425	under the Securities Act (17 CF	FR 230.425)
☐ Soliciting material pursuant to Rule 14a-12 unc	ler the Exchange Act (17 CFR 2	240.14a-12)
☐ Pre-commencement communications pursuant	to Rule 14d-2(b) under the Exc	hange Act (17 CFR 240.14d-2(b))
☐ Pre-commencement communications pursuant	to Rule 13e-4(c) under the Excl	nange 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, par value \$0.001 per share	CYCC	The Nasdaq Capital Market LLC
Preferred Stock, \$0.001 par value	CYCCP	The Nasdaq Capital Market LLC
Indicate by check mark whether the registrant is a chapter) or Rule 12b-2 of the Securities Exchange Act		defined in Rule 405 of the Securities Act of 1933 (§230.405 of this ter).
Emerging growth company \square		
If an emerging growth company, indicate by chec new or revised financial accounting standards provided		ed not to use the extended transition period for complying with any Exchange Act. \Box

Item 3.01 Notice of Delisting or Failure to Satisfy a Continued Listing Rule or Standard; Transfer of Listing.

On October 15, 2024, Cyclacel Pharmaceuticals, Inc. (the "Company") met with the Nasdaq Hearings Panel regarding the Company's potential delisting from The Nasdaq Stock Market as a result of its violation of the equity requirement in Listing Rule 5550(b)(1) (the "Equity Rule") or any of the alternative requirements in Listing Rule 5550(b). On October 22, 2024, the Company received the Nasdaq Hearings Panel decision which granted the Company until December 24, 2024 to regain compliance with the Equity Rule. If the Company is unable to regain compliance with the listing standards of the Nasdaq Capital Market by December 24, 2024, the Company's securities may be delisted from The Nasdaq Stock Market.

Item 9.01. Financial Statements and Exhibits.

(d) Exhibits

Exhibit No. Description

99.1 Press Release of Cyclacel Pharmaceuticals, Inc., dated October 24, 2024

104 Cover Page Interactive Data File (embedded with 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.

CYCLACEL PHARMACEUTICALS, INC.

By: /s/ Paul McBarron

Name: Paul McBarron

Title: Executive Vice President-Finance,

Chief Financial Officer and Chief Operating Officer

Date: October 24, 2024





PRESS RELEASE

CYCLACEL PHARMACEUTICALS ANNOUNCES THAT NASDAQ GRANTED AN EXTENSION TO REGAIN COMPLIANCE WITH THE EQUITY STANDARD RULE

BERKELEY HEIGHTS, NJ, October 24, 2024 -- Cyclacel Pharmaceuticals, Inc. (NASDAQ: CYCC, NASDAQ: CYCCP; "Cyclacel" or the "Company"), a biopharmaceutical company developing innovative cancer medicines, today announced that it received a written notice (the "Notice") from The Nasdaq Stock Market LLC ("Nasdaq") granting the Company an extension until December 24, 2024, to regain compliance with Nasdaq's minimum stockholders equity requirement under Nasdaq Listing Rule 5550(b)(1) (the "Equity Standard Rule"). This Notice has no immediate effect on the listing of the Company's common stock on Nasdaq.

As previously reported, on August 26, 2024, the Listing Qualifications Staff of Nasdaq (the "Staff") determined that the Company was not in compliance with the Equity Standard Rule because the Company reported stockholders' equity of less than \$2.5 million as of June 30, 2024.

Notwithstanding the foregoing, there can be no assurance that the Company will be able to meet the deadlines or conditions imposed by the Hearings Panel, or regain compliance with all applicable requirements for continued listing. Additionally, the Nasdaq Listing and Hearing Review Council may, on its own motion, determine to review any Hearing Panel decision within 45 calendar days after issuance of the written decision. If the Listing Council determines to review the Hearing Panel's decision, it may affirm, modify, reverse, dismiss or remand the decision to the Hearing Panel.

About Cyclacel Pharmaceuticals, Inc.

Cyclacel is a clinical-stage, biopharmaceutical company developing innovative cancer medicines based on cell cycle, transcriptional regulation and mitosis biology. The transcriptional regulation program is evaluating fadraciclib, a CDK2/9 inhibitor, and the anti-mitotic program plogosertib, a PLK1 inhibitor, in patients with both solid tumors and hematological malignancies. Cyclacel's strategy is to build a diversified biopharmaceutical business based on a pipeline of novel drug candidates addressing oncology and hematology indications. For additional information, please visit www.cyclacel.com.

Forward-looking Statements

This news release contains certain forward-looking statements that involve risks and uncertainties that could cause actual results to be materially different from historical results or from any future results expressed or implied by such forward-looking statements. Such forward-looking statements include, among other things, statements related to the Company's ability to regain compliance with Nasdaq's stockholders equity requirement and remain listed on Nasdaq. Factors that may cause actual results to differ materially include market and other conditions, the risk that product candidates that appeared promising in early research and clinical trials do not demonstrate safety and/or efficacy in larger-scale or later clinical trials, trials may have difficulty enrolling, Cyclacel may not obtain approval to market its product candidates, the risks associated with reliance on outside financing to meet capital requirements, the risks associated with reliance on collaborative partners for further clinical trials, development and commercialization of product candidates and Cyclacel's ability to regain and maintain compliance with Nasdaq's continued listing requirements. You are urged to consider statements that include the words "may," "will," "would," "could," "should," "believes," "estimates," "projects," "potential," "expects," "plans," "anticipates," "intends," "continues," "forecast," "designed," "goal," or the negative of those words or other comparable words to be uncertain and forward-looking. For a further list and description of the risks and uncertainties the Company faces, please refer to our most recent Annual Report on Form 10-K and other periodic and other filings we file with the Securities and Exchange Commission and are available at www.sec.gov. Such forward-looking statements are current only as of the date they are made, and we assume no obligation to update any forward-looking statements, whether as a result of new information, future events or otherwise.

Contact

Company: Paul McBarron, (908) 517-7330, IR@cyclacel.com

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