

**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): December 16, 2024

Vaccinex, Inc.

(Exact name of registrant as specified in its charter)

Delaware
(State or other jurisdiction
of incorporation)

001-38624
(Commission
File Number)

16-1603202
(IRS Employer
Identification No.)

1895 Mount Hope Avenue, Rochester, New York
(Address of principal executive offices)

14620
(Zip Code)

Registrant's telephone number, including area code: (585) 271-2700

(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, par value \$0.0001 per share	VCNX	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 (§ 230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§ 240.12b-2 of this chapter).

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

On December 16, 2024, Vaccinex, Inc. (the “Company”) received written notice (the “Notice”) from the Office of General Counsel of The Nasdaq Stock Market (“Nasdaq”) indicating that the Nasdaq Hearings Panel (the “Panel”) has determined to delist the Company’s shares from Nasdaq due to the Company’s failure to meet Nasdaq’s continued listing standards. As previously disclosed, the Company has not been compliant with the requirements under Nasdaq Listing Rule 5550(b)(1) to maintain a minimum of \$2.5 million in stockholders’ equity for continued listing on the Nasdaq Capital Market. The Notice indicated that trading in the Company’s shares of common stock (the “Common Stock”) on Nasdaq will be suspended effective at the open of trading on Wednesday, December 18, 2024.

Upon suspension of the trading of its Common Stock on Nasdaq, the Company expects that its Common Stock will be quoted under its existing symbol “VCNX” on the OTC Markets Group.

Item 8.01 Other Events.

On December 17, 2024, the Company issued a press release announcing, among other things, the Panel’s delisting determination. A copy of the press release is attached as Exhibit 99.1 and incorporated by reference herein.

Forward-Looking Statements

Except for the factual statements made herein, information contained in this report consists of forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995 that involve risks, uncertainties and assumptions that are difficult to predict. Words such as “will,” “may,” “intends,” “plans,” and similar expressions, or the use of future tense, identify forward-looking statements, but their absence does not mean that a statement is not forward-looking. Such forward-looking statements are not guarantees of performance and actual actions or events could differ materially from those contained in such statements. For example, there can be no assurance that the Company’s Common Stock will be quoted on the OTC Markets Group. The risks included are not exhaustive; for a more detailed description of these uncertainties and other factors, see the other risk factors set forth from time to time in the Company’s filings with the Securities and Exchange Commission, copies of which are available for free at www.sec.gov or upon request from the Company’s Investor Relations Department. All information provided in this Form 8-K is as of the date hereof and the Company assumes no obligation and does not intend to update these forward-looking statements, except as required by law.

Item 9.01 Financial Statements and Exhibits.

The following exhibits are filed herewith:

<u>Exhibit Number</u>	<u>Exhibit Description</u>
99.1	Press Release, dated December 17, 2024
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 hereunto duly authorized.

Vaccinex, Inc.

Date: December 17, 2024

By: /s/ Jill Sanchez
Jill Sanchez
Chief Financial Officer



Vaccinex Announces Receipt of Delisting Notification from Nasdaq

ROCHESTER, N.Y., December 17, 2024 (GLOBE NEWSWIRE) — Vaccinex, Inc. (Nasdaq: VCNX) (“Vaccinex” or the “Company”), a clinical-stage biotechnology company pioneering a differentiated approach to treating neurodegenerative disease by blocking astroglialosis and neuroinflammation through the inhibition of SEMA4D, today announced that on December 16, 2024, the Company received written notice (the “Notice”) from the Office of General Counsel of The Nasdaq Stock Market (“Nasdaq”) indicating that the Nasdaq Hearings Panel has determined to delist the Company’s shares from Nasdaq due to the Company’s failure to meet Nasdaq’s continued listing standards. As previously disclosed, the Company has not been compliant with the requirements under Nasdaq Listing Rule 5550(b)(1) to maintain a minimum of \$2.5 million in stockholders’ equity for continued listing on the Nasdaq Capital Market. The Notice indicated that trading in the Company’s shares of common stock (the “Common Stock”) on Nasdaq will be suspended effective at the open of trading on Wednesday, December 18, 2024.

Upon suspension of the trading of its Common Stock on Nasdaq, the Company expects that its Common Stock will be quoted under its existing symbol “VCNX” on the OTC Markets Group.

About Vaccinex, Inc.

Vaccinex, Inc. is pioneering a differentiated approach to treating slowly progressive neurodegenerative diseases and cancer through the inhibition of semaphorin 4D (SEMA4D). The Company’s lead drug candidate, pepinemab, blocks SEMA4D, a potent biological effector that it believes triggers damaging inflammation in chronic diseases of the brain and prevents infiltration and activation of immune cells in tumors. Pepinemab was studied as a monotherapy in the Phase 1b/2 SIGNAL-AD study in Alzheimer’s Disease, and the Company has previously published promising Phase 2 data in Huntington’s disease. Vaccinex believes pepinemab could also be an important contributor to combination therapy in AD. In oncology, pepinemab is being evaluated in combination with KEYTRUDA® in the Phase 1b/2 KEYNOTE-B84 study in recurrent or metastatic head and neck cancer (HNSCC) and in combination with BAVENCIO® in a Phase 1b/2 study in patients with metastatic pancreatic adenocarcinoma (PDAC). The oncology clinical program also includes several investigator-sponsored studies in solid tumors including breast cancer and melanoma.

Vaccinex has global commercial and development rights to pepinemab and is the sponsor of the KEYNOTE-B84 study which is being performed in collaboration with Merck Sharp & Dohme Corp, a subsidiary of Merck and Co, Inc. Kenilworth, NJ, USA.

KEYTRUDA is a registered trademark of Merck Sharp & Dohme Corp., a subsidiary of Merck & Co. Inc., Kenilworth, NJ, USA.

BAVENCIO®/avelumab is provided by Merck KGaA, Darmstadt, Germany, previously as part of an alliance between the healthcare business of Merck KGaA, Darmstadt, Germany and Pfizer.

About Pepinemab

Pepinemab is a humanized IgG4 monoclonal antibody designed to block SEMA4D, which can bind to plexin-B1 receptors to trigger collapse of the actin cytoskeleton in cells and lead to loss of homeostatic functions of astrocytes and other glial cells in the brain and of dendritic cells in immune tissue. Pepinemab appears to have been well-tolerated with a favorable safety profile in multiple clinical trials in different neurological and cancer indications.



Forward Looking Statements

To the extent that statements contained in this press release are not descriptions of historical facts regarding Vaccinex, Inc. (“Vaccinex,” “we,” “us,” or “our”), they may be forward-looking statements reflecting management’s current beliefs and expectations. Such statements include, but are not limited to, statements identified by words such as “may,” “will,” “intends,” “plans,” and similar expressions or their negatives (as well as other words and expressions referencing future events, conditions, or circumstances). These statements include, among others, those regarding the expected timing of the quotation of the Company’s Common Stock on the OTC Markets Group. These statements are based on our current expectations and beliefs and are subject to a number of factors and uncertainties that could cause actual results to differ materially from those described in the forward-looking statements. Except as required by law, we assume no obligation to update these forward-looking statements. For a further discussion of these and other factors that could cause future results to differ materially from any forward-looking statement, see the section titled “Risk Factors” in our periodic reports filed with the Securities and Exchange Commission (the “SEC”) and the other risks and uncertainties described in Vaccinex’s most recent year-end Annual Report on Form 10-K and subsequent SEC filings.

Investor Contact

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