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 10, 2022

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.) 14620

(Zip Code)

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

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

(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)

Dere-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 \boxtimes

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.

Vaccinex, Inc. (the "Company") will present at the Jefferies Healthcare Conference on Friday, June 10, 2022 at 10:30 a.m. Eastern Time. A copy of the corporate presentation that will accompany the presentation is furnished to this Current Report on Form 8-K as Exhibit 99.1 and is available on the Company's website located at <u>www.vaccinex.com</u> under the heading "Investors" and subheading "Presentations."

The information furnished pursuant to this Item 7.01, including Exhibit 99.1, 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 under such section and shall not be deemed to be incorporated by reference into any filing of the Company under the Securities Act of 1944, as amended, or the Exchange Act.

Item 9.01 Financial Statements and Exhibits.

Exhibit No.	Description
99.1	June 2022 corporate presentation of Vaccinex, Inc.
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: June 10, 2022

By: /s/ Scott E. Royer Scott E. Royer Chief Financial Officer

Pepinemab – Anti-SEMA4D Antibody for Cancer Immunotherapy



KEYNOTE-B84 Study in HNSCC

June 10, 2022

Forward Looking Statements

To the extent that statements contained in this presentation are not descriptions of historical facts regarding Vaccinex, Inc. ("Vaccinex," "we," "us," or "our"), they are forward-looking statements reflecting management's current beliefs and expectations. Such statements include, but are not limited to, statements about the Company's plans, expectations and objectives with respect to the results and timing of clinical trials of pepinemab in various indications, the use and potential benefits of pepinemab in Head and Neck cancer, Huntington's and Alzheimer's disease and other indications, and other statements identified by words such as "may," "will," "appears," "expect," "planned," "anticipate," "estimate," "intend," "hypothesis," "potential," "advance," and similar expressions or their negatives (as well as other words and expressions referencing future events, conditions, or circumstances). Forward-looking statements involve substantial risks and uncertainties that could cause the outcome of the Company's research and pre-clinical development programs, clinical development programs, future results, performance, or achievements to differ significantly from those expressed or implied by the forward-looking statements. Such risks and uncertainties include, among others, uncertainties inherent in the execution, cost and completion of preclinical and clinical trials, uncertainties related to regulatory approval, the risks related to the Company's dependence on its lead product candidate pepinemab, the ability to leverage its ActivMAb[®] platform, the impact of the COVID-19 pandemic, and other matters that could affect the Company's development plans or the commercial potential of its product candidates. Except as required by law, the Company assumes 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 the Company's periodic reports filed with the Securities and Exchange Commission ("SEC") and the other risks and uncertainties described in the Company's most recent year end Annual Report on Form 10-K and subsequent filings with the SEC.

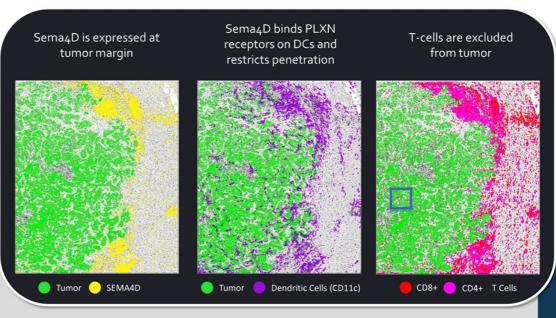




WHY DOES IMMUNE RESPONSE FAIL IN TUMORS?

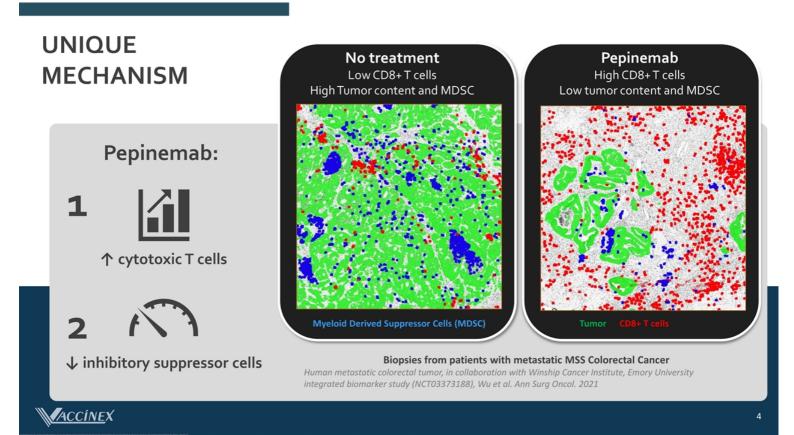
Immune Exclusion Activated T-cells and den

Activated T-cells and dendritic cells can't penetrate tumor



Pro-inflammatory cells are excluded from tumor and build up at the invasive edge CD8 T cells align with Sema4D at the invasive edge of the tumor. Most of these excluded T-cells express Sema4D. Dendritic Cells express receptors for SEMA4D and are heavily excluded at the invasive edge. Human metastatic colorectal tumor, in collaboration with Emory University (NCT03373188)

ACCINEX



RATIONALE FOR TREATMENT OF HNSCC

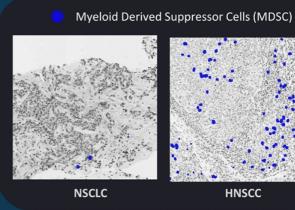
Head and Neck cancer (HNSCC)

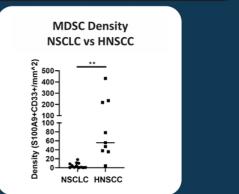
- Data suggest that SEMA4D is strongly expressed in HNSCC & induces high levels of myeloid derived suppressor cells (MDSC)
- Relatively low (17-19%) response rate to immune checkpoint therapy in HNSCC



Hypothesis: Inhibiting MDSC with pepinemab may enhance response to pembrolizumab in HNSCC







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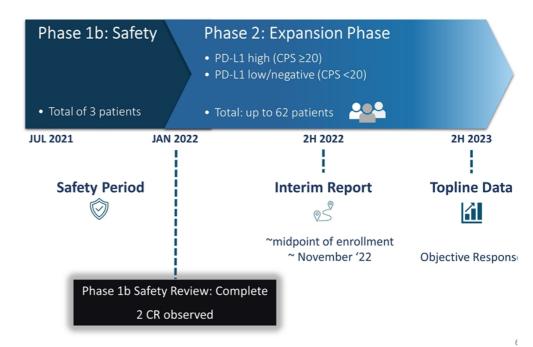
KEYNOTE B84 HEAD AND NECK CANCER TRIAL

- All patients receive standard of care Keytruda[®], plus pepinemab for firstline treatment
- Ph1b Safety: COMPLETE
 - Appeared to be well tolerated
 - RP2D: 20mg/kg pepi and 200mg pembro, Q3W
- Ph2 Expansion: Accruing
- 17 of 18 sites in USA now actively enrolling
- > Open-label, continuous monitoring



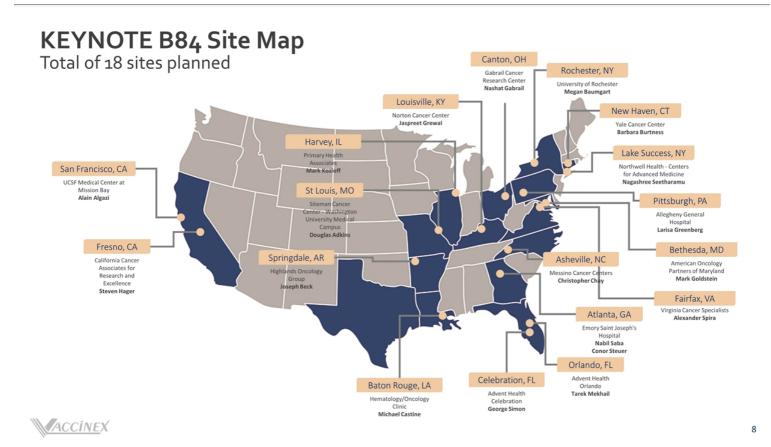
MERCK

KEYNOTE B84: pepinemab + Keytruda[®] for first-line treatment of recurrent or metastatic head and neck cancer



KEYNOTE-B84: Case studies to illustrate differences in positive response profiles

			•			
Case Study # 1: CR (confirmed)	Biopsy week 5	week 9	Scans week 15	week 21	Biomarkers	Adverse Events
Oropharyngeal cancer Target lesions: metastases to lung (Left 11mm, Right 15mm)	NO malignancy	19% decrease, SD	100% decrease, CR	Confirmed, CR Now 42 Weeks	PD-L1 CPS<20 HPV negative	none of notable severity
Case Study # 2: CR (confirmed)						
Larynx cancer with direct invasion into thyroid and neck Target lesions: neck mass (37mm)	NO malignancy	100% decrease, CR	Confirmed, CR	Continued CR Now 27 Weeks	PD-L1 CPS<1 HPV negative	Grade 1 rash
Case Study # 4: PR (unconfirmed)						
Oropharyngeal cancer Target lesions: metastases to lung (Left 24 mm, Right 23 mm)	Not available	6% decrease, SD	72% decrease, PR	Anticipated July '22	PD-L1 CPS≥20 HPV negative	none of notable severity

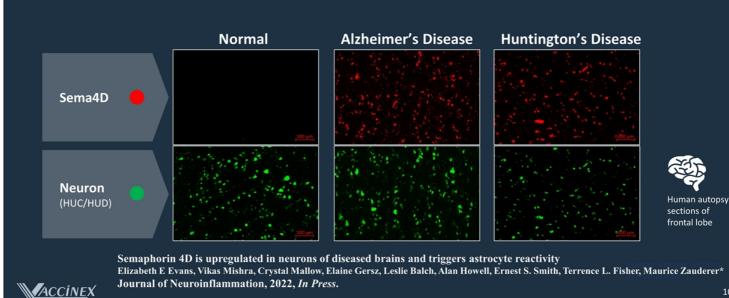


Pepinemab – Anti-SEMA4D Antibody for Huntington's and Alzheimer's Disease



Novel Mechanisms New Medicines

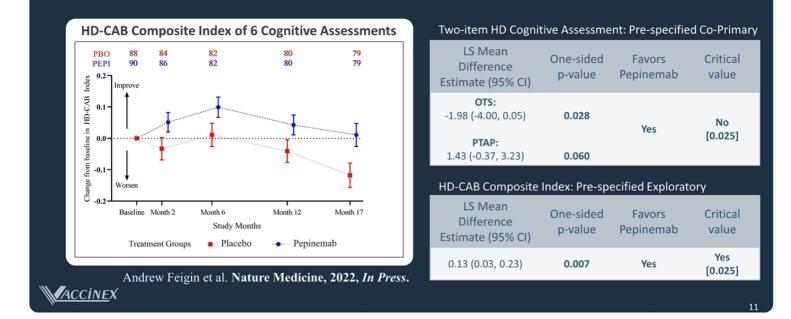
SEMA4D IS UPREGULATED IN NEURONS DURING ALZHEIMER'S AND HUNTINGTON'S DISEASE PROGRESSION



COGNITIVE ASSESSMENT BATTERY (HD-CAB)

Co-Primary and pre-specified Exploratory analysis





COGNITIVE ASSESSMENT BATTERY (HD-CAB)

Exploratory and Post-hoc analysis



18

6 12 Study month

BL

HD-CAB composite score B1 HD-CAB composite score - placebo PBO B1 87 87 84 PBO B2 43 43 42 82 42 **80** 39 "Learning effect" is lost when 87 87 84 89 89 86 82 82 80 80 0.2 0.2-▲ Change from baseline (Z-score) (Z-score) pepinemab HD symptoms become 0.1-0.1prodromal manifest 0.0 line 0.0 -0.1 -0.1 Pepinemab treatment restores 1 mon -0.2-¥ * placebo early manifest the ability to benefit from -0.2 Change p=0.007 -0.3 -0.3 experience (ie, to learn) 12 18 18 BL 6 Study month BL 6 12 Study month HD-CAB composite score, MoCA<26 HD-CAB composite score, MoCA ≥26 Pepinemab antibody blockade of SEMA4D in PBO 50 50 PEPI 53 51 PBO 36 34 PEPI 36 35 32 33 33 32 32 32 47 47 47 50 49 early Huntington's Disease: the randomized, 0.2-0.2 from baseline (Z-score) (7-score) pepinemab t 1 placebo-controlled, phase 2 SIGNAL trial 0.1 0.1-Andrew Feigin, Elizabeth E. Evans, Terrence L. Fisher, John E. Leonard, Ernest S. Smith, Alisha Reader, Vikas Mishra, haseline 0.0 0.0 Richard Manber, Kimberly A. Walters, Lisa Kowarski, David Oakes, Eric Siemers, Karl D. Kieburtz, Maurice Zauderer*, and the Huntington Study Group SIGNAL investigators **Nature Medicine**, 2022, *In Press*. -0.1 -0.1 from -0.2 -0.2 placebo Change 1 Change p=0.0025 -0.3 -0.3

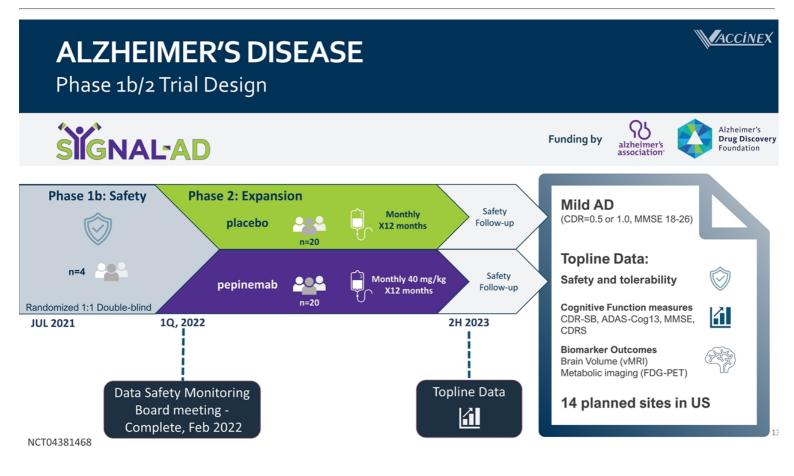
12

Study month

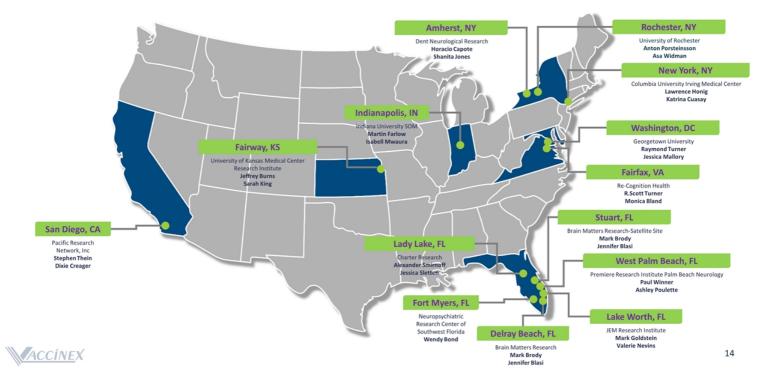
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Signal-AD Site Map





PIPELINE and MILESTONES

Research/Preclinical	Phase 1	Phase 2	Phase 3	Partner/Funding	Milestone
Pepinemab Antibody Platfo	orm (anti-Sema	ohorin 4D Mab)			
Oncology					
Pepinemab COMBO with Non Small Cell Lung Car			CLASSICAL- Lung	Merck, KGaA Darmstadt	Complete, Published 2021
Pepinemab COMBO with Head and Neck Cancer	h Pembrolizumab	in	KEYNOTE- B84	MERCK Merck, MSD	Ongoing Next data 2H 202
leurology					
Pepinemab in Huntingto (Orphan Drug and Fast Trac			SIGNAL		Complete, Nat. Med in pres
Pepinemab in Alzheime	r's Disease		SIGNAL-AD	Alzheimer's alzheimer's association: Alzheimer's Drug Discovery Foundation	Ongoing Data 2H 2023
				All Studies Sponsored by:	