

**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): August 12, 2021**

**Vincerx Pharma, Inc.**  
(Exact name of registrant as specified in its charter)

**Delaware**  
(State or Other Jurisdiction  
of Incorporation)

**001-39244**  
(Commission  
File Number)

**83-3197402**  
(I.R.S. Employer  
Identification No.)

**260 Sheridan Avenue, Suite 400**  
**Palo Alto, California**  
(Address of principal executive offices)

**94306**  
(Zip Code)

**(650) 800-6676**  
(Registrant's telephone number, including area code)

**N/A**  
(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>
<b>Common Stock, \$0.0001 par value per share</b>	<b>VINC</b>	<b>The Nasdaq Stock Market LLC</b>

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.

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**Item 2.02 Results of Operations and Financial Condition.**

On August 12, 2021, Vincerx Pharma, Inc. (the “Company”) issued a press release announcing financial results for its fiscal quarter ended June 30, 2021. The full text of the press release is furnished as Exhibit 99.1 to this Current Report on Form 8-K.

The information in Item 2.02 of this Current Report on Form 8-K, including Exhibit 99.1, shall not be deemed to be filed for purposes of Section 18 of the Securities Exchange Act of 1934 (the “Exchange Act”), or otherwise subject to the liability of that section, and shall not be incorporated by reference into any registration statement or other document filed under the Securities Act of 1933 or the Exchange Act, except as shall be expressly set forth by specific reference in such filing.

**Item 9.01 Financial Statements and Exhibits.**

(d) Exhibits.

99.1 [Press release dated August 12, 2021.](#)

104 Cover Page Interactive Data File (embedded within the Inline XBRL document)

SIGNATURE

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.

Dated: August 12, 2021

VINCERX PHARMA, INC.

By:   /s/ Raquel E. Izumi    
Raquel E. Izumi  
President and Chief Operations Officer

**Vincerx Pharma Reports Second Quarter 2021 Financial Results and Provides a Corporate Update**

*First patient dosed in Phase 1b study of VIP152 in MYC-driven relapsed or refractory aggressive lymphomas and advanced solid tumors*

*Phase 1b dose escalation study of VIP152 in patients with CLL relapsed/refractory to venetoclax and BTK inhibitors on track to initiate in 2H2021*

PALO ALTO, California, August 12, 2021 — Vincerx Pharma, Inc. (Nasdaq: VINC), a biopharmaceutical company aspiring to address the unmet medical needs of patients with cancer through paradigm-shifting therapeutics, today reported financial results for the second quarter ended June 30, 2021 and provided a corporate update.

“During the past quarter, we achieved an important milestone in dosing the first patient in our Phase 1b study of VIP152, our selective PTEFb/CDK9 inhibitor, in MYC-driven indications, building upon the compelling signals of monotherapy activity observed in the dose-escalation study and exploratory cohort in double-hit lymphoma,” said Ahmed Hamdy M.D., Chief Executive Officer of Vincerx. “Looking ahead, we intend to continue our strong execution in the clinic with the initiation of our Phase 1 dose escalation study in CLL relapsed/refractory to venetoclax and BTK inhibitors in the second half of this year. Our balance sheet provides us with a solid foundation to continue to execute on our clinical and regulatory goals.”

**Recent Highlights**

- Announced first patient dosed in Phase 1b study of VIP152, a potent and selective inhibitor of CDK9, in MYC-driven relapsed or refractory aggressive lymphomas and advanced solid tumors
  - Ongoing Phase 1b expansion, first-in-human (FIH) study is in patients with advanced cancer and consists of two expansion arms. Arm 1 will enroll up to 30 patients with relapsed/refractory aggressive lymphoma, including DLBCL, transformed follicular lymphoma, and blastoid mantle cell lymphoma. Arm 2 will enroll up to 40 patients with advanced solid tumors, including patients with ovarian cancer, triple negative breast cancer, castration-resistant neuroendocrine prostate cancer, and any other solid tumor with MYC aberration. All patients must have confirmed MYC overexpression or translocation.
- Presented clinical data in a poster presentation entitled “*Safety and efficacy of VIP152, a PTEFb / CDK9 inhibitor, in patients with double-hit lymphoma*” at the 2021 American Society of Clinical Oncology (ASCO) Annual Meeting
- Published an article entitled “*Changing for the Better: Discovery of the Highly Potent and Selective CDK9 Inhibitor VIP152 Suitable for Once Weekly Intravenous Dosing for the Treatment of Cancer,*” in the *Journal of Medicinal Chemistry*
- Announced inclusion in the Russell 3000® and Microcap® Indexes

## Second Quarter 2021 Financial Results

- Vincerx Pharma ended the second quarter with \$85.6 million in cash and cash equivalents, which includes the proceeds from the recent public warrant redemption, compared to \$61.8 million at December 31, 2020.
- Net loss for the second quarter ended June 30, 2021 was \$2.0 million, or \$0.12 per share, basic and diluted.
- Research and development (R&D) expenses were \$10.7 million for the quarter ended June 30, 2021, consisting primarily of \$3.0 million in headcount related costs, \$3.3 million of outside services in preparation for and support of our clinical trials and \$4.4 million in stock-based compensation expense.
- General and administrative (G&A) expenses were \$6.7 million for the quarter ended June 30, 2021, consisting primarily of \$1.2 million in headcount related costs, \$3.3 million of outside services in support of our operations as a public company and \$2.2 million in stock-based compensation expense.

## About Vincerx Pharma, Inc.

Vincerx Pharma, Inc. (Vincerx) is a clinical-stage life sciences company focused on leveraging its extensive development and oncology expertise to advance new therapies intended to address unmet medical needs for the treatment of cancer. Vincerx has assembled a management team of biopharmaceutical experts with extensive experience in building and operating organizations that develop and deliver innovative medicines to patients. Vincerx's current pipeline is derived from an exclusive license agreement with Bayer and includes a clinical-stage and follow-on small molecule drug program and a preclinical stage bioconjugation platform, which includes next-generation antibody-drug conjugates and innovative small molecule drug conjugates. For more information, please visit [www.vincerx.com](http://www.vincerx.com).

## Cautionary Statement

This press release contains forward-looking statements within the meaning of Section 27A of the Securities Act of 1933, as amended (the "Securities Act"), and Section 21E of the Securities Exchange Act of 1934, as amended, that are intended to be covered by the "safe harbor" created by those sections. Forward-looking statements, which are based on certain assumptions and describe future plans, strategies, expectations and events, can generally be identified by the use of forward-looking terms such as "believe," "expect," "may," "will," "should," "would," "could," "seek," "intend," "plan," "goal," "project," "estimate," "anticipate" or other comparable terms. All statements other than statements of historical facts included in this press release are forward-looking statements. Forward-looking statements include, but are not limited to: Vincerx's business model, pipeline, strategy, timeline, product candidates, preclinical and clinical development and results and future capital requirements. Forward-looking statements are neither historical facts nor assurances of future performance or events. Instead, they are based only on current beliefs, expectations and assumptions regarding future business developments, future plans and strategies, projections, anticipated events and trends, the economy and other future conditions. Forward-looking statements are subject to inherent uncertainties, risks and changes in circumstances that are difficult to predict and many of which are outside of our control.

Actual results, conditions and events may differ materially from those indicated in the forward-looking statements. Therefore, you should not rely on any of these forward-looking statements. Important factors that could cause actual results, conditions and events to differ materially from those indicated in the forward-looking statements include, but are not limited to: general economic, financial, legal, political and business conditions and changes in domestic and foreign markets; the potential effects of the COVID-19 pandemic; risks associated with preclinical or clinical development conducted prior to Vincerx's in-licensing; failure to realize the anticipated benefits of the business combination with LifeSci Acquisition Corp.; failure to realize the benefits of Vincerx's license agreement with Bayer; risks related to the rollout of Vincerx's business and the timing of expected business milestones; changes in the assumptions underlying Vincerx's expectations regarding its future business or business model; Vincerx's ability to develop and commercialize product candidates; the availability and uses of capital; the effects of competition on Vincerx's future business; and the risks and uncertainties set forth in Forms 10-K, 10-Q and 8-K filed with or furnished to the SEC from time to time by Vincerx. Forward-looking statements speak only as of the date hereof, and Vincerx disclaims any obligation to update any forward-looking statements.

**Contact Information**

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**Vincerox Pharma, Inc.**  
**Condensed Consolidated Balance Sheets**  
(in thousands)

	June 30, 2021 (Unaudited)	December 31, 2020
<b>ASSETS</b>		
Current assets:		
Cash and cash equivalents	\$ 85,623	\$ 61,792
Prepaid expenses	882	1,104
Other current assets	—	214
Total current assets	86,505	63,110
Right-of-use assets	4,217	—
Other assets	415	82
<b>Total assets</b>	<b>\$ 91,137</b>	<b>\$ 63,192</b>
<b>LIABILITIES AND STOCKHOLDERS' EQUITY</b>		
Current liabilities		
Accounts payable	1,830	505
Accrued expenses	2,397	—
Lease liability	360	—
License payable	—	5,000
Common stock warrant liabilities	11,097	32,308
Total current liabilities	15,684	37,813
Lease liability, net of current portion	3,858	—
<b>Total liabilities</b>	<b>19,542</b>	<b>37,813</b>
Total stockholders' equity	71,595	25,379
<b>Total liabilities and stockholders' equity</b>	<b>\$ 91,137</b>	<b>\$ 63,192</b>

**Vincerx Pharma, Inc.**  
**Condensed Consolidated Statements of Operations**  
**(unaudited)**  
**(in thousands, except per share amounts)**

	For the three months ended June 30,		For the six months ended June 30,	
	2021	2020	2021	2020
<b>Operating expenses:</b>				
General and administrative	\$ 6,695	\$ 33	\$ 11,486	\$ 37
Research and development	10,698	—	15,532	—
Total operating expenses	<u>17,393</u>	<u>33</u>	<u>27,018</u>	<u>37</u>
<b>Loss from operations</b>	<u>(17,393)</u>	<u>(33)</u>	<u>(27,018)</u>	<u>(37)</u>
<b>Other income</b>				
Change in fair value of warrant liabilities	15,359	—	18,708	—
Total other income	<u>15,359</u>	<u>—</u>	<u>18,708</u>	<u>—</u>
<b>Net loss</b>	<u>\$ (2,034)</u>	<u>\$ (33)</u>	<u>\$ (8,310)</u>	<u>\$ (37)</u>
Net loss per common share, basic and diluted	<u>\$ (0.12)</u>	<u>\$ (0.01)</u>	<u>\$ (0.55)</u>	<u>\$ (0.01)</u>
Weighted average common shares outstanding, basic and diluted	<u>16,350</u>	<u>5,009</u>	<u>15,050</u>	<u>4,988</u>