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): May 10, 2021

KRYSTAL BIOTECH, INC.

(Exact name of registrant as specified in its charter)

Delaware (State or other jurisdiction of incorporation) 001-38210 (Commission File Number) 82-1080209 (IRS Employer Identification Number)

2100 Wharton Street, Suite 701
Pittsburgh, Pennsylvania 15203
(Address of principal executive offices, including Zip Code)

Registrant's telephone number, including area code: (412) 586-5830

Check the a following p	11 1	ng is intended to simultaneously satisfy the filing o	obligation of the registrant under any of the					
	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	KRYS	Nasdaq					

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. \boxtimes

Item 2.02 Results of Operation and Financial Condition.

On May 10, 2021, Krystal Biotech, Inc., a Delaware corporation (the "Company"), announced its financial results for the quarter ending March 31, 2021. A copy of the Company's press release is attached as Exhibit 99.1 hereto and incorporated by reference herein.

The information concerning financial results in this Form 8-K and in Exhibit 99.1 shall not be deemed to be "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended, or otherwise subject to the liabilities of that section. The information concerning financial results in this Form 8-K and in Exhibit 99.1 shall not be incorporated into any registration statement or other document filed with the Securities and Exchange Commission by the Company, whether made before or after the date hereof, regardless of any general incorporation language in such filing, except as shall be expressly set forth by specific reference in such filing.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits.

Exhibit No.	<u>Description</u>
99.1	Press Release, dated May 10, 2021
104	Cover Page Interactive Data File (embedded within the Inline XBRL document).

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, as amended, the registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

Date: May 10, 2021 KRYSTAL BIOTECH, INC.

By: /s/ Krish S. Krishnan

Name: Krish S. Krishnan

Title: Chairman and Chief Executive Officer



Krystal Biotech Reports First Quarter 2021 Financial Results and Provides Update on Operational Progress

- · Completed enrollment in pivotal GEM-3 study of B-VEC in dystrophic epidermolysis bullosa (DEB)
- · Strong balance sheet with March 31, 2021 cash, cash equivalents and short-term investments of \$403.4 million

PITTSBURGH, May 10, 2021 – <u>Krystal Biotech Inc.</u>, ("Krystal") (NASDAQ: KRYS), the leader in redosable gene therapies for rare diseases, today reported financial results and key operational progress updates for the first quarter ending March 31, 2021.

"2021 has already been, and will continue to be, a busy period for Krystal and I would like to thank our employees for their commitment, passion and resiliency during a global pandemic, "said Krish Krishnan, Chairman and CEO of Krystal Biotech, Inc. "We look forward to continued progress across the organization, including initiating clinical testing of our first respiratory program, making headway on our 2nd GMP manufacturing facility, and announcing top line pivotal data for B-VEC later this year."

Program Highlights & Upcoming Events:

B-VEC for DEB

- During the first quarter, the Company <u>completed enrollment</u> in the ongoing pivotal GEM-3 trial and <u>finalized</u> the statistical analysis plan (SAP) based on feedback from the FDA.
- Topline data from the study is anticipated in 4Q21.
- Details of the pivotal Phase 3 study can be found at www.clinicaltrials.gov under NCT identifier NCT04491604.

KB105 for TGM1-ARCI

• The Company is on track to provide an update on this program in 1H21. The update will consist of safety, TGM1 expression level and efficacy data from the 4th patient enrolled. Data from this patient, together with the data from the three initial patients will help determine next steps.



Details of the Phase 1/2 study can be found at www.clinicaltrials.gov under NCT identifier NCT04047732.

KB407 for Cystic Fibrosis

- In April 2021, the Company <u>announced</u> data from the GLP toxicology and biodistribution study which demonstrated in vivo safety of repeat, nebulized doses of KB407 in nonhuman primates. More detailed data will be presented at a future scientific conference.
- The Company intends to initiate a Phase 1 study of KB407 in 3Q21.

KB408 for Alpha-1 Antitrypsin Deficiency

• In April 2021, Krystal <u>announced</u> initial positive proof-of-concept data from preclinical studies of KB408 in mice. Additional preclinical studies are ongoing, and more detailed data will be presented at a future scientific conference.

KB301 for Aesthetic Indications

- In May 2021, detailed safety data from the first cohort in the ongoing Phase 1 study of KB301 for aesthetic skin indications was presented at the Society of Investigative Dermatology (SID) Annual Meeting.
- Initial efficacy data from the Phase 1 study is anticipated in 2H21.

KB104 for Netherton Syndrome

• The Company continues to work towards an IND filing, which is anticipated in 2H21.

Corporate Highlights:

- In May 2021, the Company <u>announced</u> the appointment of Andy Orth as Chief Commercial Officer. Mr. Orth has more than 25 years of global biotechnology experience, most recently in the role of Senior Vice President, Head of US Business at Alnylam Pharmaceuticals.
- In March 2021, the Company <u>announced</u> the appointment of Dr. Bhuhsan Hardas, M.D., MBA as President of Jeune, Inc. the Company's wholly owned subsidiary that is leveraging Krystal's clinically validated gene delivery platform to fundamentally address and reverse the biology of aging and/or damaged skin. Before joining Jeune, Dr. Hardas served as Chief Scientific Officer, Executive Vice President, Global Head of Licensing at Almirall and previously served as Chief Medical Officer of Allergan's Dermatology and Medical Aesthetics business.



- In February 2021, the Company <u>completed</u> a public offering of 2,211,538 shares of its common stock, including 288,461 shares purchased by the underwriters, at \$65.00 per share. Net proceeds to the Company from the offering were \$134.9 million.
- In January 2021, 262,500 shares of common stock were issued pursuant to our at-the-market equity offering program ("ATM Program") for net proceeds of \$16.9 million, resulting in a remaining \$132.5 million available for issuance under the ATM Program. The Company also incurred \$172 thousand of other offering expenses related to the ATM Program.

Financial results for the quarter ended March 31, 2021:

- Cash, cash equivalents and short-term investments totaled \$403.4 million on March 31, 2021.
- Research and development expenses for the first quarter ended March 31, 2021 were \$6.2 million, compared to \$3.5 million for the first quarter 2020.
- General and administrative expenses for the first quarter ended March 31, 2021 were \$8.2 million, compared to \$2.4 million for first quarter 2020.
- Net losses for the quarters ended March 31, 2021 and 2020 were \$15.8 million and \$5.3 million, or \$(0.74) and \$(0.31) respectively, per common share (basic and diluted).
- For additional information on the Company's financial results for the first quarter ended March 31, 2021, refer to form 10-Q filed with the SEC.

About Krystal Biotech

Krystal Biotech, Inc. (NASDAQ:KRYS) is a pivotal-stage gene therapy company leveraging its novel, redosable gene therapy platform and in-house manufacturing capabilities to develop therapies to treat serious rare diseases. For more information please visit http://www.krystalbio.com.

Forward-Looking Statements

Any statements in this press release about future expectations, plans and prospects for Krystal Biotech, Inc., including but not limited to statements about the development of Krystal's product candidates, such as plans for the design, conduct and timelines of ongoing pre-clinical and clinical trials of beremagene geperpavec ("B-VEC"), KB105, KB104, KB301, KB407, and KB408; the clinical utility of B-VEC, KB105, KB104, KB301, KB407 and KB408, and Krystal's plans for filing of regulatory approvals and efforts to bring B-VEC, KB105, KB104, KB301, KB407 and



KB408 to market; the market opportunity for and the potential market acceptance of B-VEC, KB105, KB104, KB301, KB407 and KB408; plans to pursue research and development of other product candidates; the sufficiency of Krystal's existing cash resources; the unanticipated impact of COVID-19 on Krystal's business operations, pre-clinical activities and clinical trials; and other statements containing the words "anticipate," "believe," "estimate," "expect," "intend," "may," "plan," "predict," "project," "target," "potential," "likely," "will," "would," "could," "should," "continue," and similar expressions, constitute forward-looking statements within the meaning of The Private Securities Litigation Reform Act of 1995. Actual results may differ materially from those indicated by such forward-looking statements as a result of various important factors, including: the uncertainties inherent in the initiation and conduct of clinical trials, availability and timing of data from clinical trials, whether results of early clinical trials or trials will be indicative of the results of ongoing or future trials, uncertainties associated with regulatory review of clinical trials and applications for marketing approvals, the availability or commercial potential of product candidates including B-VEC, KB105, KB104, KB301, KB407 and KB408, the sufficiency of cash resources and need for additional financing and such other important factors as are set forth under the caption "Risk Factors" in Krystal's annual and quarterly reports on file with the U.S. Securities and Exchange Commission. In addition, the forward-looking statements included in this press release represent Krystal's views as of the date of this release. Krystal anticipates that subsequent events and developments will cause its views to change. However, while Krystal may elect to update these forward-looking statements at some point in the future, it specifically disclaims any obligation to do so. These forward-looking statements should not be relied upon as representi

CONTACTS:

Investors:

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Consolidated Balance Sheet Data:

(In thousands)		December 31, 2020
Balance sheet data:		
Cash and cash equivalents	\$402,172	\$ 268,269
Working capital	396,681	259,606
Total assets	\$443,018	\$ 310,844
Total liabilities	12,366	18,760
Total stockholders' equity	\$430,652	\$ 292,084

Consolidated Statement of Operations:

		Three Months Ended March 31,			
(In thousands, except shares and per share data)		2021		2020	Change
Expenses					
Research and development	\$	6,201	\$	3,525	\$ 2,676
General and administrative		8,152		2,421	5,731
Total operating expenses		14,353		5,946	8,407
Loss from operations		(14,353)		(5,946)	(8,407)
Other Income (Expense)					
Interest and other income, net		33		605	(572)
Build to suit interest expense		(1,492)			(1,492)
Total other income (expense)		(1,459)		605	(2,064)
Net loss		(15,812)		(5,341)	(10,471)
Net loss applicable to stockholders		(15,812)	\$	(5,341)	\$(10,471)
Net loss attributable to common stockholders per share: Basic and diluted		(0.74)	\$	(0.31)	
Weighted-average common shares outstanding: Basic and diluted		1,253,508	17	7,359,356	

