UNITED STATES SECURITIES AND EXCHANGE COMMISSION

		WASHINGTON, D.C. 20347	
		FORM 8-K	
		CURRENT REPORT	
	Pursuant to Sect	ion 13 or 15(d) of the Securities Exchange	Act of 1934
	Date of I	Report (Date of earliest event reported): May 13, 2	.022
		Nkarta, Inc. (Exact name of Registrant as Specified in Its Charter)	
	Delaware (State or Other Jurisdiction of Incorporation)	001-39370 (Commission File Number)	47-4515206 (IRS Employer Identification No.)
	6000 Shoreline Court, Suite 102 South San Francisco, CA (Address of Principal Executive Offices)		94080 (Zip Code)
	Registrant's	Telephone Number, Including Area Code: (415) 5	82-4923
	(Fo	Not Applicable rmer Name or Former Address, if Changed Since Last Report)	
	ck the appropriate box below if the Form 8-K filir owing provisions:	ng is intended to simultaneously satisfy the filing obli	gation of the registrant under any of the
	Written communications pursuant to Rule 425 u	under the Securities Act (17 CFR 230.425)	
П	Soliciting material pursuant to Rule 14a-12 and	er the Evchange Act (17 CER 240 14a-12)	

			(Nasdaq Global Select Market)				
Common Stock, \$0.0001 par value per share		NKTX	The Nasdaq Stock Market LLC				
	Title of each class	Trading Symbol(s)	Name of each exchange on which registered				
Secu	Securities registered pursuant to Section 12(b) of the Act:						
	Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))						
	Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))						
	Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)						

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 5.02 Departure of Directors or Certain Officers; Election of Directors; Appointment of Certain Officers; Compensatory Arrangements of Certain Officers.

On May 13, 2022, Dr. Kanya Rajangam notified Nkarta, Inc. (the "Company") of her decision to resign as Chief Medical Officer of the Company to oversee research and development activities at a private biotechnology company. Dr. Rajangam's resignation is effective June 5, 2022.

Item. 7.01 Regulation FD Disclosure.

On May 16, 2022, the Company issued a press release announcing the resignation of Dr. Rajangam and the appointment of Dr. David Shook as Vice President, Clinical Development. A copy of the press release is attached hereto as Exhibit 99.1 and incorporated herein by reference.

The information in Item 7.01 of this Current Report on Form 8-K (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 of that section, nor shall it be, or be deemed, incorporated by reference in any filings under the Securities Act of 1933, as amended (the "Securities Act"), unless the Company specifically states that the information is to be considered "filed" under the Exchange Act or incorporates it by reference into a filing under the Securities Act or the Exchange Act.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits.

Exhibit Number	Description		
99.1	Press Release issued on May 16, 2022.		
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 regis hereunto duly authorized.	trant has duly caused this report to be signed on its behalf by the undersigned		
	Nkarta, Inc.		
Date: May 16, 2022	By: /s/ Alicia Hager Alicia J. Hager, J.D., Ph.D. Chief Legal Officer		



Nkarta Appoints David R. Shook, MD, as Vice President, Clinical Development

SOUTH SAN FRANCISCO, Calif., May 16, 2022 -- Nkarta, Inc. (Nasdaq: NKTX), a clinical-stage biopharmaceutical company developing engineered natural killer (NK) cell therapies to treat cancer, today announced the appointment of David R. Shook, MD, as Vice President, Clinical Development. Dr. Shook is a practicing pediatric hematologist, oncologist, and transplanter, and an early pioneer of natural killer (NK) cell therapy. He currently directs Nkarta's co-lead clinical programs, NKX101 and NKX019. In his expanded role, Dr. Shook will lead all clinical development and regulatory activities at Nkarta. Kanya Rajangam, MD, PhD, has resigned as Chief Medical Officer, effective June 5, 2022, and has accepted a position to oversee research and development activities at a private biotechnology company.

"On behalf of Nkarta's Board of Directors, I wish to thank Kanya for her many contributions to the company's progress. Under her leadership, NKX101 and NKX019 reached crucial early development milestones and demonstrated preliminary evidence of anti-tumor activity in AML and NHL, respectively," said Paul J. Hastings, President and CEO of Nkarta. "We are excited to welcome David, a seasoned and valued member of Nkarta's clinical team, to his new position. This will be a seamless transition given David's experience in leading the ongoing development of our co-lead programs, and his foundational work with Nkarta's scientific founder, Dr. Dario Campana, on the engineering and enhancement of NK cells. We look forward to leveraging David's considerable clinical expertise and understanding of Nkarta's technology to advance our cell therapy candidates."

Nkarta's clinical programs continue to progress. In April 2022, Nkarta reported preliminary single-agent proof of concept data for its independent co-lead clinical programs, NKX101 and NKX019. As previously announced, Nkarta is currently evaluating a higher-dose regimen of 1.5 billion x 3 doses of CAR NK cells in the dose escalation portion of the NKX101 and NKX019 Phase 1 clinical studies. Nkarta expects to submit updated data from these studies for presentation at a medical meeting this year.

About David R. Shook, MD

Dr. Shook has more than 10 years of clinical research and development experience. Prior to joining Nkarta in June 2020, he led multiple first-in-human cell therapy clinical trials, including CD19 CAR-NK and CD45RA-depleted BMT. He was a fellow, fellowship director and faculty member at St. Jude Children's Research Hospital, where he conducted research in the laboratory of Dario Campana, PhD, Nkarta's scientific founder. Dr. Campana and Dr. Shook co-discovered the membrane bound form of interleukin-15 (IL-15), a key component of Nkarta's engineered NK cell platform technology. Dr. Shook has authored dozens of scientific publications covering areas including the quantitation of minimal residual disease in acute myeloid leukemia, pediatric stem cell transplantation, and NK cell and T cell therapies for hematologic malignancies and solid tumors. He is board certified in Pediatric Hematology & Oncology and General Pediatrics. He earned his medical degree from The Johns Hopkins University School of Medicine and his bachelor's degree from Purdue University.

About NKX101

NKX101 is an allogeneic, cryopreserved, off-the-shelf cancer immunotherapy candidate that uses natural killer (NK) cells derived from the peripheral blood of healthy donors. It is engineered with a chimeric antigen

receptor (CAR) targeting NKG2D ligands on tumor cells. NKG2D, a key activating receptor found on naturally occurring NK cells, induces a cell-killing immune response through the detection of stress ligands that are widely expressed on cancer cells. NKX101 is also engineered with membrane-bound form of interleukin-15 (IL15) for greater persistence and activity without exogenous cytokine support. To learn more about the NKX101 clinical trial in adults with AML or MDS, please visit <u>ClinicalTrials.gov</u>.

About NKX019

NKX019 is an allogeneic, cryopreserved, off-the-shelf cancer immunotherapy candidate that uses natural killer (NK) cells derived from the peripheral blood of healthy adult donors. It is engineered with a humanized CD19-directed CAR for enhanced tumor cell targeting and a proprietary, membrane-bound form of interleukin-15 (IL-15) for greater persistence and activity without exogenous cytokine support. CD19 is a biomarker for normal and malignant B cells, and it is a validated target for B cell cancer therapies. To learn more about the NKX019 clinical trial in adults with advanced B cell malignancies, please visit ClinicalTrials.gov.

About Nkarta

Nkarta is a clinical-stage biotechnology company advancing the development of allogeneic, off-the-shelf natural killer (NK) cell therapies for cancer patients. By combining its cell expansion and cryopreservation platform with proprietary cell engineering technologies and CRISPR-based genome engineering capabilities, Nkarta is building a pipeline of future cell therapies engineered for deep anti-tumor activity and intended for broad access in the outpatient treatment setting. For more information, please visit the company's website at www.nkartatx.com.

Cautionary Note on Forward-Looking Statements

Statements contained in this press release regarding matters that are not historical facts are "forward-looking statements" within the meaning of the Private Securities Litigation Reform Act of 1995, as amended. Words such as "anticipates," "believes," "expects," "intends," "plans," "potential," "projects," "would," and "future" or similar expressions are intended to identify forward-looking statements. Examples of these forward-looking statements include statements concerning Nkarta's expectations regarding any or all of the following: Nkarta's ability to continue to build and advance its pipeline of clinical and preclinical product candidates; the potential impact of changes in Nkarta's leadership; the timing of release of additional NKX019 and NKX101 clinical trial data; the anti-tumor activity and safety profile of NKX019 and NKX101; and the ability of Nkarta's technology to augment the anti-tumor activity of NK cells and enable broad access. Interim clinical data referenced in this press release were reported on April 25, 2022 and are subject to the risk that one or more of the clinical outcomes may materially change as patient enrollment continues and more data on existing patients become available.

Because forward looking statements are subject to risks and uncertainties, actual results may differ materially from those expressed or implied by such forward-looking statements. These risks and uncertainties include, among others: Nkarta's limited operating history and historical losses; Nkarta's lack of any products approved for sale and its ability to achieve profitability; the risk that the results of preclinical studies and early-stage clinical trials may not be predictive of future results; Nkarta's ability to raise additional funding to complete the development and any commercialization of its product candidates; Nkarta's dependence on the success of its co-lead product candidates, NKX101 and NKX019; that Nkarta may be delayed in initiating, enrolling or completing any clinical trials; competition from third parties that are developing products for similar uses; Nkarta's ability to obtain, maintain and protect its intellectual property; Nkarta's dependence on third parties in connection with manufacturing, clinical trials, and pre-clinical studies; the complexity of the manufacturing process for CAR NK cell therapies; and risks relating to the impact on Nkarta's business of the COVID-19 pandemic or similar public health crises.

These and other risks are described more fully in Nkarta's filings with the Securities and Exchange Commission ("SEC"), including the "Risk Factors" section of Nkarta's Quarterly Report on Form 10-Q for the quarter ended March 31, 2022, filed with the SEC on May 12, 2022, and Nkarta's other documents subsequently filed with or furnished to the SEC. All forward-looking statements contained in this press release speak only as of the date on which they were made. Except to the extent required by law, Nkarta undertakes no obligation to update such statements to reflect events that occur or circumstances that exist after the date on which they were made.

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