### 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 2, 2022 (July 29, 2022)

# **C4 THERAPEUTICS, INC.**

(Exact name of Registrant as Specified in Its Charter)

Delaware (State or Other Jurisdiction of Incorporation) 001-39567

(Commission File Number)

490 Arsenal Way, Suite 120 Watertown, MA (Address of Principal Executive Offices) Identification No.) 02472

47-5617627

(IRS Employer

(Zip Code)

Registrant's Telephone Number, Including Area Code: (617) 231-0700

Not Applicable

(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, \$0.0001 par value per share	CCCC	The Nasdaq Global Select 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  $\Box$ 

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.  $\Box$ 

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

On July 29, 2022, the Company's Board of Directors (the "**Board**"), upon the recommendation of the Nominating and Corporate Governance Committee of the Board, elected Donna Grogan, M.D. and Laura Bessen, M.D, as directors of the Company.

Dr. Bessen has been designated as a Class I director, to serve until the Company's 2024 Annual Meeting of Stockholders, and Dr. Grogan has been designated as a Class II director, to serve until the Company's 2025 Annual Meeting of Stockholders, in accordance with the Company's Second Amended and Restated By-Laws and each to serve thereafter until her successor has been duly elected and qualified or until her earlier death, removal or resignation.

In connection with these elections, the Company granted to each of Dr. Bessen and Dr. Grogan stock options to purchase up to 41,200 shares of the Company's common stock under the Company's 2020 Stock Option and Incentive Plan. One-third of each of these stock option awards shall vest on the first anniversary of the date of grant, with the remainder vesting quarterly over the subsequent two years, *provided*, *however*, that all vesting of each award shall cease if its recipient resigns from the Board or otherwise ceases to serve as a director of the Company or otherwise maintain a service relationship with the Company prior to any such vesting date. In addition, in connection with these elections, the Company entered into an indemnification agreement with each of Dr. Bessen and Dr. Grogan in the same form as used with the Company's other directors.

There are no arrangements or understandings between either of Dr. Bessen or Dr. Grogan and any other persons pursuant to which she was selected as a director of the Company, and there are no transactions in which either of Dr. Bessen or Dr. Grogan has an interest requiring disclosure under Item 404(a) of Regulation S-K.

#### Item 7.01 Regulation FD Disclosure.

On August 2, 2022, the Company issued a press release announcing the elections of Dr. Bessen and Dr. Grogan as directors. A copy of the press release is attached as Exhibit 99.1 to this Current Report on Form 8-K.

The information in Item 7.01 of this Current Report on Form 8-K, including Exhibit 99.1, is being furnished and shall not be deemed "filed" for the 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 deemed subject to the requirements of amended Item 10 of Regulation S-K, nor shall it be deemed incorporated by reference into any filing of the Company under the Securities Act of 1933, as amended, or the Exchange Act, whether made before or after the date hereof, regardless of any general incorporation language in such filing. The furnishing of this information hereby shall not be deemed an admission as to the materiality of any such information.

#### Item 9.01 Financial Statements and Exhibits.

(d) Exhibits. The exhibits shall be deemed to be filed or furnished, depending on the relevant item requiring such exhibit, in accordance with the provisions of Item 601 of Regulation S-K (17 CFR 229.601) and Instruction B.2 to this form.

Exhibit Number	Description
99.1	Press release issued August 2, 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 registrant has duly caused this report to be signed on its behalf by the undersigned thereunto duly authorized.

C4 Therapeutics, Inc.

Date: August 2, 2022

By: /s/ Jolie M. Siegel

Jolie M. Siegel Chief Legal Officer



# C4 Therapeutics Appoints Experienced Clinical Development Leaders Laura Bessen, M.D. and Donna Grogan, M.D. to Board of Directors

Dr. Laura Bessen Brings More Than Two Decades of Experience Across Medical Affairs and Clinical Development in Support of Successful Product Launches –

 Dr. Donna Grogan Joins as Accomplished Drug Development and Regulatory Strategy Leader With More than 25 Years of Experience in Developing Novel Therapeutics –

Appointments Highlight C4T's Continued Commitment to Evolving Board Composition –

WATERTOWN, Mass., August 2, 2022 (GLOBE NEWSWIRE) – C4 Therapeutics, Inc. (C4T) (Nasdaq: CCCC), a clinical-stage biopharmaceutical company dedicated to advancing targeted protein degradation science to develop a new generation of small-molecule medicines and transform how disease is treated, today announced the appointments of Laura Bessen, M.D. and Donna Grogan, M.D. to its board of directors.

"We are delighted to announce Laura and Donna as C4T's newest directors after an extensive search that included a talented and diverse pool of clinical development leaders experienced in helping companies bring innovative therapies to patients," said Bruce Downey, chairman of C4 Therapeutics. "On behalf of the board of directors, I look forward to working closely with Laura and Donna as C4T continues to evolve in our pursuit of delivering on the potential of targeted protein degradation science to create a new generation of medicines and change how diseases are treated."

"I am eager to use my broad experiences in medical affairs and clinical development to help C4T advance multiple oncology programs to benefit patients," said Laura Bessen, M.D. "C4T is at a transformational stage, with two programs in the clinic and additional programs preparing to advance into clinical trials. I am excited to help C4T capitalize upon opportunities ahead."

Commenting on joining the C4T board of directors, Donna Grogan, M.D. said, "It is an exciting time to join C4T and partner with the company to navigate drug development and regulatory milestones to advance an innovative portfolio of small-molecule medicines. I look forward to working with my C4T colleagues to help the Company apply its industry-leading targeted protein degradation science to transform how patients are treated."

#### About Laura Bessen, M.D.



Laura Bessen, M.D. currently serves as managing partner at Maxsam Advisors LLC where she provides strategic clinical and medical affairs advice to clients across the biotechnology and pharmaceutical industries. Previously, she held roles of increasing responsibility over a 15-year career at Bristol Myers Squibb (BMS), most recently as Vice President, Head of US Medical. During her tenure at BMS, she helped launch 11 new products including Opdivo<sup>®</sup>, Yervoy<sup>®</sup>, elotuzomab, and Eliquis<sup>®</sup> and co-led BMS' partnership with Gilead to develop Atripla<sup>®</sup>. Dr. Bessen also co-led the development of BMS product commercialization

model,

life cycle management and launch investment principles. Earlier in her career, Dr. Bessen served in medical affairs roles at DuPont Pharmaceuticals. She currently serves on the Board of Directors of Artiva Biotherapeutics, an oncology company developing and advancing off-the-shelf, allogeneic natural killer cell therapies for patients with hematologic cancers or solid tumors. Dr. Bessen received her M.D. degree from New York University School of Medicine and B.S. in biochemistry from the State University of New York at Binghamton.

#### About Donna Grogan, M.D.



Donna Grogan, M.D. currently serves as Principal of Grogan Consulting LLC where she supports clients across drug development, regulatory strategy, trial design and data interpretation. Between September 2013 and June 2019, Dr. Grogan served as Chief Medical Officer of Clementia Pharmaceuticals, which was acquired by Ipsen in April 2019. She previously served as Chief Medical Officer for several HealthCare Ventures portfolio companies including Anexon, Apofore, and DecImmune. Between February 2007 and ugust 2011, Dr. Grogan served as Chief Medical Officer, Senior Vice President Clinical Development at

Pharmaceuticals which was acquired by Pfizer in October 2010. Earlier in her career, she held roles of increasing responsibility at Sepracor, Inc. where she was involved in multiple high-profile product approvals including Lunesta<sup>®</sup>, Xopenex HFA<sup>®</sup>, and Brovana<sup>TM</sup>. Dr. Grogan previously served as a board member, including membership on the Compensation Committee and the Scientific Committee, of Momenta Pharmaceuticals until it was acquired by J&J in October 2020. She holds a M.D. from University of Illinois College of Medicine and a B.A. from College of the Holy Cross.

#### **About C4 Therapeutics**

C4 Therapeutics (C4T) (Nasdaq: CCCC) is a clinical-stage biopharmaceutical company dedicated to delivering on the promise of targeted protein degradation science to create a new generation of medicines that transforms patients' lives. C4T is leveraging its TORPEDO® platform to efficiently design and optimize small-molecule medicines that harness the body's natural protein recycling system to rapidly degrade disease-causing proteins, offering the potential to overcome drug resistance, drug undruggable targets and improve patient outcomes. C4T is advancing multiple targeted oncology programs to the clinic and expanding its research platform to deliver the next wave of medicines for difficult-to-treat diseases. For more information, please visit www.c4therapeutics.com.

#### **Forward-Looking Statements**

This press release contains "forward-looking statements" of C4 Therapeutics, Inc. within the meaning of the Private Securities Litigation Reform Act of 1995. These forward-looking statements may include, but may not be limited to, express or implied statements regarding our ability to develop potential therapies for patients; the design and potential efficacy of our therapeutic approaches; the predictive capability of our TORPEDO® platform in the development of novel, selective, orally bioavailable degraders; the potential timing, design and advancement of our pre-clinical studies and clinical trials, including the potential timing for regulatory authorization related to clinical trials; our ability and the potential to successfully manufacture and supply our product candidates for clinical trials; our ability to replicate results achieved in our pre-clinical studies or clinical trials in any future studies or trials; and regulatory developments in the United States and foreign countries. Any forward-looking statements in this press release are based on management's current expectations and beliefs of future events and are subject to a number of risks and uncertainties that could cause actual results to differ materially and adversely from those set forth in or implied by such forward-looking statements. These risks and uncertainties include, but are not limited to: uncertainties related to the initiation, timing, advancement and conduct of pre-clinical and clinical studies and other development requirements for our product candidates will cost more to develop or may not be successfully developed and commercialized; and the risk that the results of pre-clinical studies and/or clinical trials will or will not be predictive of results in

<sup>,</sup> FoldRx

connection with future studies or trials. For a discussion of these and other risks and uncertainties, and other important factors, any of which could cause our actual results to differ from those contained in the forward-looking statements, see the section entitled "Risk Factors" in C4 Therapeutics' most recentAnnual Report on Form 10-K and/or Quarterly Report on Form 10-Q, as filed with the Securities and Exchange Commission. All information in this press release is as of the date of the release, and C4 Therapeutics undertakes no duty to update this information unless required by law.

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