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): July 6, 2020

CORBUS PHARMACEUTICALS HOLDINGS, INC.

(Exact name of registrant as specified in its charter)

Delaware	**001-37348**	**46-4348039**
(State or other jurisdiction of incorporation)	*(Commission File Number)*	*(IRS Employer Identification No.)*

500 River Ridge Drive, Norwood, MA	**02062**
(Address of principal executive offices)	*(Zip Code)*

Registrant's telephone number, including area code: **(617) 963-0100**

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 (see General Instruction A.2. below):

[ ] 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))

Title of each class	Trading Symbol(s)	Name of each exchange on which registered
Common Stock	CRBP	The Nasdaq Global 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 [ ]

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 July 6, 2020, the Board of Directors (the "Board") of Corbus Pharmaceuticals Holdings, Inc. (the "Company"), upon the recommendation of the Nominating and Corporate Governance Committee of the Board, appointed George S. Golumbeski, Ph.D., age 63, to serve as a member of the Board. Dr. Golumbeski will hold this position until the next annual meeting of the Company's stockholders or until his successor is elected and qualified, subject to his earlier resignation or removal.

Dr. Golumbeski is an independent biotechnology advisor since April 2018. From 2017 until April 2018, Dr. Golumbeski was the Executive Vice President for Innovation for Celgene Corporation, a biotechnology company, where he was previously Executive Vice President of Business Development since 2009, responsible for the full array of business development activities and focused primarily within the therapeutic areas of oncology and inflammation. From 2008 to 2009, Dr. Golumbeski served as the CEO of Nabriva Therapeutics AG. Prior to Nabriva, Dr. Golumbeski served as Vice President of Business Development, Licensing and Strategy for Novartis-Oncology. Earlier in his career, Dr. Golumbeski held senior positions at Elan Pharmaceuticals and at Schwarz Pharma. He currently serves on the boards of directors of Enanta Pharmaceuticals, Inc., MorphoSys AG and Sage Therapeutics, public biotechnology companies. Dr. Golumbeski received a BA in biology from the University of Virginia and a Ph.D. in genetics from the University of Wisconsin-Madison.

Dr. Golumbeski will participate in the Company's standard non-employee director compensation plan, including an initial option grant to purchase 81,000 shares of the Company's common stock, par value $0.0001 per share ("Common Stock") upon joining the Board, an annual cash retainer fee of $40,000 (pro-rated for the current year), and an annual stock option grant to purchase shares of the Company's Common Stock.

There are no transactions between Dr. Golumbeski and the Company that would be reportable under Item 404(a) of Regulation S-K.

Concurrently with the appointment, the Company entered into an indemnification agreement with Dr. Golumbeski (the "Indemnification Agreement"), in the form previously entered into by the Company with each of the Company's directors and executive officers, the form of which was filed as Exhibit 10.15 to the Amendment No. 1 to Company's Registration Statement on Form S-1 filed with the Securities and Exchange Commission on September 30, 2014. The Indemnification Agreement, subject to limitations contained therein, will obligate the Company to indemnify Dr. Golumbeski, to the fullest extent permitted by applicable law, for certain expenses, including attorneys' fees, judgments, penalties, fines and settlement amounts actually and reasonably incurred by him in any threatened, pending or completed action, suit, claim, investigation, inquiry, administrative hearing, arbitration or other proceeding arising out of his services as a director. Subject to certain limitations, the Indemnification Agreement provides for the advancement of expenses incurred by the indemnitee, and the repayment to the Company of the amounts advanced to the extent that it is ultimately determined that the indemnitee is not entitled to be indemnified by the Company. The Indemnification Agreement also creates certain rights in favor of the Company, including the right to assume the defense of claims and to consent to settlements. The Indemnification Agreement does not exclude any other rights to indemnification or advancement of expenses to which the indemnitee may be entitled under applicable law, the certificate of incorporation or bylaws of the Company, any agreement, a vote of stockholders or disinterested directors, or otherwise.

The foregoing is a summary of the material terms of the Indemnification Agreement and does not purport to be complete.

Item 7.01. Regulation FD Disclosure.

On July 8, 2020, the Company issued a press release announcing the appointment of Dr. Golumbeski to its Board of Directors. A copy of the press release is furnished as Exhibit 99.1 hereto and shall not be deemed "filed" for the purposes of Section 18 of the Exchange Act, or otherwise subject to the liabilities of that section, nor shall it be deemed incorporated by reference in any filing under the Exchange Act or the Securities Act of 1933, as amended, except as shall be expressly set forth by specific reference in such a filing.

Item 9.01. Financial Statements and Exhibits.

(d) The following exhibit is furnished with this report:

Exhibit No.	Description
99.1	Press Release of the Company dated July 8, 2020.

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.

CORBUS PHARMACEUTICALS HOLDINGS, INC.

Dated: July 8, 2020

By: */s/ Yuval Cohen*
Name: Yuval Cohen
Title: Chief Executive Officer

Exhibit 99.1

Corbus Pharmaceuticals Appoints George Golumbeski, Ph.D., to Board of Directors

- *Dr. Golumbeski formerly led corporate and business development at Celgene and Novartis*
- *Appointment adds unique skill set to Corbus Board ahead of several key data readouts this summer*

Norwood, MA, [July 8, 2020]—Corbus Pharmaceuticals Holdings, Inc. (NASDAQ: CRBP) ("Corbus" or the "Company"), a clinical-stage drug development company pioneering transformative medicines that target the endocannabinoid system (ECS), today announced the appointment of George Golumbeski, Ph.D., to its Board of Directors.

During his corporate career, Dr. Golumbeski held senior leadership positions in business development at Celgene, Novartis, Elan Pharmaceuticals and Schwarz Pharma. In these roles, he drove business strategies that created value, fostered internal and external collaboration, and built creative deal structures and teams to grow those businesses. At Celgene, he served as Executive Vice President and was responsible for the entire array of business development activities (R&D collaborations, Licensing Agreements, M&A and Alliance Management). During his nine years at the Company, he and his colleagues established Celgene as "Partner of Choice" in the biotechnology industry. Dr. Golumbeski currently serves as a Director on the Boards of Enanta Pharmaceuticals, MorphoSys, Sage Therapeutics as well as several privately held companies.

Dr. Golumbeski commented, "I am really pleased to join the team at Corbus. Philosophically, I am highly aligned with Corbus' focus on non-incremental scientific innovation directed at the highest degree of unmet medical needs. Lenabasum is a first-in-class molecule, and systemic sclerosis, as well as dermatomyositis, currently have no approved therapies for the overall treatment of disease. This makes the work underway at Corbus both very exciting and very important. I look forward to working with the Board and the Executive Team as Corbus approaches many strategic and corporate development milestones."

"The breadth, depth and sustained excellence of George's experience and accomplishments make him an ideal Board member at this pivotal time for the Company. We are honored he has chosen to join us," said Alan Holmer, Chairman of Corbus' Board of Directors.

"George's decades of experience growing companies and advancing innovation will be a significant asset to Corbus as we transition from a pre-commercial R&D organization to a commercial company with a deep pipeline of novel drug candidates, all targeting the ECS," stated Yuval Cohen, Ph.D., Chief Executive Officer of Corbus. "With George's addition to the board, the potential for what we can do with our ECS platform is greatly enhanced."

About Lenabasum

Lenabasum is a rationally designed, oral, small molecule that selectively binds as an agonist to the cannabinoid receptor type 2 (CB2), resolves inflammation, and limits fibrosis. CB2 is preferentially expressed on activated immune cells and on fibroblasts, muscle cells, and endothelial cells. In both animal and human studies conducted to date, lenabasum has induced the production of pro-resolving lipid mediators that activate endogenous pathways which resolve inflammation and speed bacterial clearance without immunosuppression. Data from animal models and human clinical studies suggest that lenabasum can reduce expression of genes and proteins involved in inflammation and fibrosis. Lenabasum has demonstrated promising activity in animal models of skin and lung inflammation and fibrosis in systemic sclerosis (SSc). Lenabasum is also active in animal models of lung infection and inflammation in cystic fibrosis and joint inflammation and scarring in rheumatoid arthritis.

Lenabasum has demonstrated acceptable safety and tolerability profiles in clinical studies to date. Lenabasum treatment was associated with improvement in multiple physician-assessed and patient-reported efficacy outcomes in Phase 2 studies in patients with diffuse cutaneous SSc and patients with dermatomyositis with active skin involvement but not currently active muscle involvement. Lenabasum treatment also was associated with a lower rate of and longer time to pulmonary exacerbations in a Phase 2 cystic fibrosis study.

Lenabasum is not approved for the treatment of systemic sclerosis, dermatomyositis, cystic fibrosis or systemic lupus erythematosus.

About Corbus

Corbus Pharmaceuticals Holdings, Inc. is a Phase 3 clinical-stage pharmaceutical company focused on the development and commercialization of novel therapeutics to treat inflammatory and fibrotic diseases by leveraging its pipeline of rationally designed, endocannabinoid system-targeting drug candidates. The Company's lead product candidate, lenabasum, is a novel, oral, selective cannabinoid receptor type 2 (CB2) agonist rationally designed to resolve chronic inflammation and fibrotic processes. Lenabasum is currently being evaluated in systemic sclerosis, cystic fibrosis, dermatomyositis and systemic lupus erythematosus.

Corbus is also developing a pipeline of drug candidates targeting the endocannabinoid system. The pipeline includes CRB-4001, a 2nd generation, selective cannabinoid receptor type 1 (CB1) inverse agonist designed to be peripherally restricted. Potential indications for CRB-4001 include nonalcoholic steatohepatitis (NASH), among others. Corbus expects data from its Phase 1 safety study in 2020.

Lenabasum is not approved for the treatment of systemic sclerosis, dermatomyositis, cystic fibrosis or systemic lupus erythematosus. CRB-4001 is not approved for the treatment of NASH/NAFLD. For more information on Corbus' clinical programs, please visit here.

Please visit www.CorbusPharma.com and connect with the Company on Twitter, LinkedIn, and Facebook.

Forward-Looking Statements

This press release contains certain forward-looking statements within the meaning of Section 27A of the Securities Act of 1933 and Section 21E of the Securities Exchange Act of 1934 and Private Securities Litigation Reform Act, as amended, including those relating to the Company's product development, clinical and regulatory timelines, market opportunity, competitive position, possible or assumed future results of operations, business strategies, potential growth opportunities and other statement that are predictive in nature. These forward-looking statements are based on current expectations, estimates, forecasts and projections about the industry and markets in which we operate and management's current beliefs and assumptions.

These statements may be identified by the use of forward-looking expressions, including, but not limited to, "expect," "anticipate," "intend," "plan," "believe," "estimate," "potential, "predict," "project," "should," "would" and similar expressions and the negatives of those terms. These statements relate to future events or our financial performance and involve known and unknown risks, uncertainties, and other factors, including the potential impact of the recent COVID-19 pandemic and the potential impact of sustained social distancing efforts, on our operations, clinical development plans and timelines, which may cause actual results, performance or achievements to be materially different from any future results, performance or achievements expressed or implied by the forward-looking statements. Such factors include those set forth in the Company's filings with the Securities and Exchange Commission. Prospective investors are cautioned not to place undue reliance on such forward-looking statements, which speak only as of the date of this press release. The Company undertakes no obligation to publicly update any forward-looking statement, whether as a result of new information, future events or otherwise.

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