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): April 26, 2019

EVOLUS, INC.

(Exact name of registrant as specified in its charter)

Delaware	001-38381	46-1385614
(State or other jurisdiction of incorporation)	(Commission File Number)	(I.R.S. Employer Identification No.)

520 Newport Center Drive, Suite 1200
Newport Beach, California 92660
(Address of principal executive offices) (Zip Code)

(949) 284-4555
(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))

Indicate by check mark whether the registrant is an emerging growth company as defined in 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 8.01 Other Events

On April 26, 2019, Evolus Inc. (the "Company") announced that the Committee for Medicinal Products for Human Use ("CHMP") of the European Medicines Agency ("EMA") adopted a positive opinion related to its Marketing Authorization Application ("MAA") during the April 2019 meeting of the CHMP for the Company's product candidate, Nuceiva™, an injectable 900 kilodalton botulinum toxin type A complex.

On April 26, 2019, the Company issued a press release announcing the information set forth above. A copy of the Company's press release is attached hereto as Exhibit 99.1 and is incorporated herein by reference.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits.

Exhibit Number	Description
99.1	Press Release of Evolus, Inc., dated April 26, 2019

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.

Evolus, Inc.

Dated: April 26, 2019

/s/ David Moatazedi

David Moatazedi

President and Chief Executive Officer

Evolus Receives Positive CHMP Opinion for Nuceiva™ in the European Union

European Commission Approval Anticipated Within 90 Days

Newport Beach, Calif., April 26, 2019 - Evolus, Inc. (NASDAQ: EOLS), a performance beauty company with a customer-centric approach focused on delivering breakthrough products, today announced the European Medicines Agency's (EMA) Committee for Medicinal Products for Human Use (CHMP) has issued a positive opinion on the Nuceiva™ marketing authorization application.

The CHMP opinion is a scientific recommendation for marketing authorization to the European Commission, which will now review the recommendation and deliver its final decision on the Company's marketing authorization application. The decision will be applicable to all 28 European Union member states plus Iceland, Norway and Liechtenstein.

"We are pleased to receive a positive CHMP opinion and look forward to receiving final expected approval of our marketing authorization application," said Rui Avelar, MD, Chief Medical Officer and Head of Research & Development.

Nuceiva™ (branded Jeuveau™ in the United States) is a proprietary 900 kDa purified botulinum toxin type A formulation for the temporary improvement in the appearance of moderate to severe vertical lines between the eyebrows seen at maximum frown (glabellar lines), when the severity of the above facial lines has an important psychological impact in adults below 65 years of age.

"The receipt of the CHMP positive opinion further validates the rigor of our *TRANSPARENCY* clinical development program, which includes the largest aesthetics head-to-head pivotal study versus BOTOX®. In anticipation of approval of Nuceiva™, we continue to evaluate commercial partnership opportunities in Europe," said David Moatazedi, President and Chief Executive Officer. "We have now achieved success with our regulatory filings for Jeuveau™ / Nuceiva™ in the United States, Canada and the European Union."

About Nuceiva™ (branded Jeuveau™ in the United States)
Nuceiva™ (prabotulinumtoxinA-xvfs) is a proprietary 900 kDa purified botulinum toxin type A formulation. Nuceiva™ is produced under strict quality and safety standards in a state-of-the art facility, specifically built to manufacture Nuceiva™. The safety and efficacy of Nuceiva™ has been evaluated in clinical studies with over 2,100 patients enrolled. Nuceiva™ is being evaluated by the EMA for the temporary improvement in the appearance of moderate to severe vertical lines between the eyebrows seen at maximum frown (glabellar lines), when the severity of the above facial lines has an important psychological impact in adults below 65 years of age.

About Evolus, Inc.

Evolus is a performance beauty company with a customer-centric approach focused on delivering breakthrough products. In 2019, the U.S. Food and Drug Administration approved Jeuveau™ (prabotulinumtoxinA-xvfs), the first and only neurotoxin dedicated exclusively to aesthetics and manufactured in a state-of-the-art facility using Hi-Pure™ technology. Jeuveau™ is powered by Evolus' unique technology platform and is designed to transform the aesthetic market by eliminating the friction points existing for customers today. Visit us at: www.evolus.com.

Forward-Looking Statements

Statements made in this press release that relate to future plans, events, prospects or performance are forward-looking statements as defined under the Private Securities Litigation Reform Act of 1995. All statements, other than statements of historical fact, are statements that could be deemed forward-looking statements, including statements containing the words "planned," "expect," "believes," "strategy," "opportunity," "anticipates," "outlook," "designed," and similar words. While these forward-looking statements are based on the current expectations and beliefs of management, such forward-looking statements

are subject to a number of risks, uncertainties, assumptions and other factors that could cause actual results to differ materially from the expectations expressed in this press release, including the risks and uncertainties disclosed in Evolus' periodic filings with the Securities and Exchange Commission, including factors described in the section entitled "Risk Factors" in its Annual Report on Form 10-K for the year ended December 31, 2018 as filed with the Securities and Exchange Commission on March 20, 2019, respectively, all of which are available online at www.sec.gov. Readers are cautioned not to place undue reliance on these forward-looking statements, which speak only as of the date hereof. Except as required by law, Evolus undertakes no obligation to update or revise any forward-looking statements to reflect new information, changed circumstances or unanticipated events.

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