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**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, 2018

**EVOLUS, INC.**

(Exact name of registrant as specified in its charter)

**Delaware**  
(State or other jurisdiction  
of incorporation)

**001-38381**  
(Commission File Number)

**46-1385614**  
(I.R.S. Employer  
Identification No.)

**17901 Von Karman Avenue, Suite 150  
Irvine, California 92614**  
(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.

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**Item 2.02 Results of Operations and Financial Condition.**

On August 2, 2018, Evolus, Inc. (the "Company") issued a press release announcing its financial results for the three and six months ended June 30, 2018. The press release is being furnished as Exhibit 99.1 to this Current Report on Form 8-K.

As provided in General Instruction B.2 of Form 8-K, the information in 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 incorporated by reference into any registration statement or other document filed under the Securities Act of 1933, as amended, or the Exchange Act, except as shall be expressly set forth by specific reference in such filing.

**Item 8.01 Other Events.**

On August 2, 2018, the Company issued a press release announcing early resubmission to the U.S. Food and Drug Administration (the "FDA") of the Company's Biologics License Application for its product candidate, PrabotulinumtoxinA (DWP-450), an injectable 900 kilodalton botulinum toxin type A complex. A copy of the press release is included as Exhibit 99.2 to this Current Report on Form 8-K.

**Item 9.01 Financial Statements and Exhibits.**

(d) Exhibits.

**Exhibit  
Number**

**Description**

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[99.1](#) [Press Release of Evolus, Inc., dated August 2, 2018](#)

[99.2](#) [Press Release of Evolus, Inc., dated August 2, 2018](#)

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**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: August 2, 2018

/s/ David Moatazedi

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David Moatazedi

President and Chief Executive Officer



## **Evolus Reports Second Quarter 2018 Financial Results**

**Irvine, Calif., August 2, 2018** – Evolus, Inc. (NASDAQ: EOLS), a company dedicated to aesthetic medicine, today reported financial results for the second quarter ended June 30, 2018.

### **Second Quarter 2018 and Recent Highlights:**

- Announced early resubmission to the U.S. Food and Drug Administration (“FDA”) of its Biologics License Application (“BLA”) for its product candidate, DWP-450.
- Closed out inspections of its partner’s DWP-450 manufacturing facility by the FDA and health agencies in the European Union and Canada.
- Appointed Lauren Silvernail as Chief Financial Officer and EVP Corporate Development
- Named Mike Jafar as Chief Marketing Officer.
- Raised net proceeds to the company of approximately \$56.4 million, before offering expenses, in a follow-on public offering of its common stock.
- Remain on track for anticipated U.S. commercial launch of DWP-450 in spring 2019.

David Moatazedi, President and Chief Executive Officer of Evolus, stated, “In the second quarter we made significant progress on key initiatives to prepare for the U.S. commercial launch of DWP-450. Most significantly, today we announced the early resubmission of our BLA. We continue to fill key positions to enable us to refine and advance our commercial strategy. Notably, we appointed Mike Jafar as Chief Marketing Officer and since his appointment have made considerable progress planning our commercial launch.”

Lauren Silvernail, Chief Financial Officer and EVP Corporate Development, added, “Through the second quarter, we continued to manage our expenses tightly, consuming only \$6.0 million of cash since March 31, 2018. In July, we completed a public equity offering, which increased our cash by \$56.4 million before offering expenses. We now believe we are well funded to take us through our early commercial stage and look forward to deploying this capital for a robust product launch.”

### **Second Quarter 2018 Financial Results**

Operating expenses for the second quarter ended June 30, 2018 were \$16.1 million, as compared to \$2.3 million in the second quarter ended June 30, 2017. The increase was primarily attributable to expenses related to the revaluation of future contingent royalties on sales of \$8.2 million, stock-based compensation of \$2.6 million which increased largely due to changes in the senior executive team, and increases in costs related to operating as a public company, offset by a reduction in clinical trial costs associated with completion of Evolus' Phase III clinical trials last year.

Non-GAAP operating expense for the second quarter ended June 30, 2018 was \$5.3 million and is calculated as operating expense, excluding stock-based compensation of \$2.6 million and a revaluation expense of \$8.2 million related to future royalties on sales.

Net loss for the second quarter ended June 30, 2018 was \$16.4 million, or \$0.69 basic and diluted net loss per share, compared with a net loss of \$2.3 million, or \$0.14 basic and diluted net loss per share, for the second quarter ended June 30, 2017.

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Total cash was \$43.6 million as of June 30, 2018, a decrease of \$6.0 million compared to March 31, 2018. Cash as of June 30, 2018 does not include total net proceeds of \$56.4 million, before offering expenses, from Evolus' follow-on common stock offering completed in July 2018.

### **Conference Call Information**

Management will host a conference call and webcast to discuss Evolus' financial results today at 8:30 a.m. ET. The dial-in numbers are (866) 916-2317 for domestic callers and (703) 925-2662 for international callers, and the conference ID is 9738939.

A replay of the call will be available following its completion through August 9, 2018. To access the replay, dial (855) 859-2056 for domestic callers and (404) 537-3406 for international callers and use the replay conference ID 9738939.

A live audio webcast of the call will be available on the Investor Relations page of the Evolus, Inc. website, <https://investors.evolus.com>. A replay of the webcast will be archived on Evolus' website for 30 days following the completion of the call.

### **About Evolus, Inc.**

Evolus is a company dedicated to aesthetic medicine focused on providing physicians and their patients with expanded choices in aesthetic treatments and procedures. Evolus' lead candidate DWP-450, also known by the chemical name prabotulinumtoxinA, is a 900 kDa purified botulinum toxin type A complex that is being evaluated for the treatment of moderate to severe glabellar lines in adult patients.

EVOLUS™ and the Evolus logo are trademarks of Evolus, Inc.

### **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, 2017; its Quarterly Report on Form 10-Q for the Quarter ended March 31, 2018; and its final prospectus for its offering, as filed with the Securities and Exchange Commission on March 29, 2018, May 10, 2018 and July 20, 2018, respectively, all of which are available online at [www.sec.gov](http://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.

### **Use of Non-GAAP Financial Measures**

Evolus' financial results are prepared in accordance with Generally Accepted Accounting Principles (“GAAP”). This press release and the reconciliation tables included in the financial schedules below include non-GAAP operating expense which is calculated as total operating expenses, excluding: (i) the revaluation of contingent royalty obligation payable and (ii) stock-based compensation expense. Management believes that non-GAAP operating expense is useful in helping to identify recurring operation performance and enables management to consistently analyze the period-to-period financial performance of the core business operations. Management also believes that non-GAAP operating expense will enable investors to assess in the same way management

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assesses Evolus' current and future operations. Non-GAAP operating expense should be considered in addition to results prepared in accordance with GAAP, but should not be considered a substitute for or superior to GAAP results.

For a reconciliation of non-GAAP Operating Expense to total operating expenses, the most directly comparable GAAP financial measure, please see “Reconciliation of GAAP Operating Expense to Non-GAAP Operating Expense” in the financial schedules below.

**Evolus Contacts:**

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**Evolus, Inc.**  
**Condensed Statements of Operations and Comprehensive Loss**  
(in thousands, except net loss per share data)  
(Unaudited)

	Three Months Ended June 30,		Six Months Ended June 30,	
	2018	2017	2018	2017
<b>Operating expenses:</b>				
Research and development	\$ 1,648	\$ 1,365	\$ 3,326	\$ 4,016
General and administrative	6,248	803	9,715	2,018
Revaluation of contingent royalty obligation payable	8,200	—	9,100	—
Depreciation and amortization	4	107	4	218
Total operating expenses	16,100	2,275	22,145	6,252
Loss from operations	(16,100)	(2,275)	(22,145)	(6,252)
<b>Other expense:</b>				
Interest expense, net	321	1	428	2
Loss before taxes	(16,421)	(2,276)	(22,573)	(6,254)
Income tax expense	12	20	22	40
Net loss and comprehensive loss	\$ (16,433)	\$ (2,296)	\$ (22,595)	\$ (6,294)
Net loss per share, basic and diluted	\$ (0.69)	\$ (0.14)	\$ (1.03)	\$ (0.38)
Weighted-average shares outstanding used to compute basic and diluted net loss per share	23,688	16,527	21,962	16,527

**Evolus, Inc.**  
**Summary of Balance Sheet Data**  
(in thousands)

	June 30, 2018 (unaudited)	December 31, 2017
<b>Balance Sheet Data:</b>		
Cash	\$ 43,585	\$ —
Working capital	40,939	(1,237)
Total assets	121,934	152,233
Total current liabilities	3,684	212,748
Total liabilities	83,999	227,776
Accumulated deficit	(98,753)	(75,543)
Total stockholders' equity (deficit)	\$ 37,935	\$ (75,543)

**Evolus, Inc.**  
**Reconciliation of GAAP Operating Expense to Non-GAAP Operating Expense**  
(in thousands)

	Three Months Ended June 30,		Six Months Ended June 30,	
	2018	2017	2018	2017
Operating expense:				
<b>GAAP operating expense</b>	\$ 16,100	\$ 2,275	\$ 22,145	\$ 6,252
Adjustments:				
Revaluation of contingent royalty obligation payable	8,200	—	9,100	—
Stock-based compensation	2,623	104	3,630	246
<b>Non-GAAP operating expense</b>	<u>\$ 5,277</u>	<u>\$ 2,171</u>	<u>\$ 9,415</u>	<u>\$ 6,006</u>



## **Evolus Announces Early Resubmission to the FDA of its Biologics License Application for DWP-450**

*BLA Resubmission to be Formally Filed Today*

*Commercial Launch Planned for Spring 2019*

**Irvine, Calif., August 2, 2018** - Evolus, Inc. (NASDAQ: EOLS), a company dedicated to aesthetic medicine, announced that today it is resubmitting its Biologics License Application (“BLA”) for its lead product candidate, DWP-450 (prabotulinumtoxinA), to the U.S Food and Drug Administration (“FDA”).

The resubmission follows the receipt of a Complete Response Letter (“CRL”) from the FDA in May 2018 which necessitated the submission of additional data to the FDA for the completion of review of Evolus’ BLA. Deficiencies cited by the FDA in the CRL were isolated to items related to Chemistry, Manufacturing, and Controls (“CMC”) processes. No deficiencies were related to clinical, non-clinical or safety matters.

David Moatazedi, President and Chief Executive Officer of Evolus, commented, “Today we are resubmitting our BLA well ahead of our guided timeline and less than 90 days since receiving our CRL. We believe this submission gives us a line of sight to the anticipated approval and subsequent commercialization of DWP-450.”

Mr. Moatazedi continued, “We look forward to receiving notice next month of the FDA’s acceptance of our resubmission and the assignment of a new FDA action date. We are working towards the planned commercial launch of DWP-450 in spring 2019 as we further accelerate the build out of our sales and marketing team and finalize our go-to-market strategy.”

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