



Evolus Announces Publication of Largest Aesthetic Head-to-Head Pivotal Trial Comparing Jeuveau™ and BOTOX® in Aesthetic Surgery Journal

April 8, 2019

Primary Endpoint of Non-Inferiority Achieved with Responder Rates of 87.2% for Jeuveau™, 82.8% for BOTOX® and 4.2% for Placebo at Day 30*

Secondary and Exploratory Analyses Unveil Additional Information about Jeuveau™ Clinical Performance

Thirty New Exploratory Head-to-Head Data Points

Conference Call and Live Webcast Scheduled for April 8th, 2019 at 8:00am ET / 5:00am PT

NEWPORT BEACH, Calif., April 08, 2019 (GLOBE NEWSWIRE) -- Evolus, Inc. (NASDAQ: EOLS), a performance beauty company with a customer-centric approach focused on delivering breakthrough products, today announced the publication of the European and Canadian Phase III head-to-head pivotal trial results in the April issue of [Aesthetic Surgery Journal](#).

"We are pleased that these data were accepted and published in the *Aesthetic Surgery Journal*, which is added validation of the rigor of the study," said Rui Avelar, MD, Chief Medical Officer and Head of Research & Development. "We knew it was crucial to invest in this study to answer key questions about the clinical performance of these two 900 kDa neurotoxins, and now clinicians can review it and draw their own conclusions."

"The results of this study are compelling and reinforce the clinical performance of Jeuveau™, which is the first 900 kDa product to enter the U.S. market in nearly 30 years," said David Moatazedi, President and Chief Executive Officer. "We look forward to launching Jeuveau™ in the coming weeks in the United States."

Conference Call and Webcast

Management will host a conference call and live webcast on Monday, April 8th, 2019 at 8:00am ET to discuss the publication. A question and answer session will follow management remarks. The dial-in numbers for the conference call are (866) 916-2317 for domestic callers and (703) 925-2662 for international callers. The conference ID is 2872109. A replay of the call will be available following its completion through April 15th, 2019. To access the replay, dial (855) 859-2056 for domestic callers and (404) 537-3406 for international callers and use the replay conference ID 2872109.

A live webcast of the conference call with accompanying slides can be accessed from the Events & Presentations page in the Investors section of Evolus' website at www.evolus.com. A replay of the webcast will be archived on Evolus' website for 30 days following the completion of the call.

About the EVB-003 Study

EVB-003 was a 150-day, multicenter, randomized, double-blind, active- and placebo-controlled, single-dose Phase III non-inferiority study evaluating prabotulinumtoxinA and onabotulinumtoxinA, both 900 kDa botulinum toxin type A complexes, in subjects who felt their glabellar lines had an important psychological impact. Adults aged 18 or older with moderate to severe glabellar lines at maximum frown, as assessed by the investigator on the 4-point Glabellar Line Scale (GLS, 0=no lines, 1=mild, 2=moderate, 3=severe), who met these criteria were enrolled. Randomization was 5:5:1 to receive a single treatment of 20 U prabotulinumtoxinA, 20 U onabotulinumtoxinA or placebo (0.9% saline).

About Jeuveau™

Jeuveau™ (prabotulinumtoxinA-xvfs) is a proprietary 900 kDa purified botulinum toxin type A formulation indicated for the temporary improvement in the appearance of moderate to severe glabellar lines in adults. Jeuveau™ is produced under strict quality and safety standards in a state-of-the-art facility, specifically built to manufacture Jeuveau™. The safety and efficacy of Jeuveau™ has been evaluated in clinical studies with over 2,100 patients enrolled.

IMPORTANT SAFETY INFORMATION FOR JEUVEAU™ (prabotulinumtoxinA-xvfs)

JEUVEAU may cause serious side effects that can be life threatening. Get medical help right away if you have any of these problems any time (hours to weeks) after injection of JEUVEAU:

- **Problems swallowing, speaking, or breathing**, due to weakening of associated muscles, can be severe and result in loss of life. You are at the highest risk if these problems are pre-existing before injection. Swallowing problems may last for several months.
- **Spread of toxin effects**. The effect of botulinum toxin may affect areas away from the injection site and cause serious symptoms including: loss of strength and all-over muscle weakness, double vision, blurred vision and drooping eyelids, hoarseness or change or loss of voice, trouble saying words clearly, loss of bladder control, trouble breathing, trouble swallowing.

Do not use JEUVEAU if you: are allergic to any of the ingredients in JEUVEAU (see Medication Guide for ingredients); had an allergic reaction to any other botulinum toxin product such as rimabotulinumtoxinB (MYOBLOC®), onabotulinumtoxinA (BOTOX®/BOTOX® Cosmetic), abobotulinumtoxinA (DYSPORE®), or incobotulinumtoxinA (XEOMIN®); have a skin infection at the planned injection site; or are a child.

Jeuveau dosing units are not the same as, or comparable to, any other botulinum.

Tell your healthcare provider about all your muscle or nerve conditions, such as ALS or Lou Gehrig's disease, Myasthenia gravis, or

Lambert-Eaton syndrome, as you may be at increased risk of serious side effects including difficulty swallowing and difficulty breathing from typical doses of JEUVEAU.

Tell your healthcare provider about all your medical conditions, including: any side effects from botulinum toxin products, including dry eye; breathing, swallowing, bleeding, or heart problems; plans to have surgery; weakness of forehead muscles; drooping eyelids; had surgery on your face; are pregnant or breastfeeding or plan to become pregnant or breastfeed (it is not known if JEUVEAU can harm your unborn baby or passes into breast milk).

Tell your healthcare provider about all the medicines you take, including prescription and over-the-counter medicines, vitamins, and herbal supplements. Using JEUVEAU with certain other medicines may cause serious side effects. **Do not start any new medicines until you have told your healthcare provider that you have received JEUVEAU in the past.**

Especially tell your healthcare provider if you: have received any other botulinum toxin product in the past and the last 4 months. and exactly which product you received (such as Botox, Botox Cosmetic, Myobloc, Dysport, or Xeomin).

JEUVEAU may cause loss of strength or general muscle weakness, vision problems, or dizziness within hours to weeks of treatment with JEUVEAU. **If this happens, do not drive a car, operate machinery, or do other dangerous activities.**

JEUVEAU can cause other serious side effects including: allergic reactions (such as itching, rash, red itchy welts, wheezing, asthma symptoms, or dizziness or feeling faint), heart problems (such as irregular heartbeat and heart attack), and eye problems (including dry eye, reduced blinking, and corneal problems). Tell your healthcare provider or get medical emergency help right away if you experience a serious side effect.

The most common side effects include: headache; eyelid drooping, upper respiratory tract infection, and increased white blood cell count in your blood.

APPROVED USE

JEUVEAU is a prescription medicine that is injected into muscles and used in adults for a short period of time (temporary) to improve the look of moderate to severe frown lines between the eyebrows (glabellar lines).

The risk information provided here is not complete. For more information about **JEUVEAU**, the full Prescribing Information including BOXED WARNING, and Medication Guide, visit www.evolus.com or talk to your healthcare provider.

To report side effects associated with use of JEUVEAU, please call 1-877-386-5871. You are encouraged to report negative side effects of prescription drugs to the FDA. Visit www.fda.gov/medwatch or call 1-800-FDA-1088.

Manufactured by: Evolus, Inc., 1027 Garden St., Santa Barbara, CA 93101

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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.

JEUVEAU™ is a trademark of Evolus, Inc.

Hi-Pure™ is a trademark of Daewoong Pharmaceutical Co, Ltd.

Botox®, Botox® Cosmetic, Myobloc®, Dysport®, and Xeomin® are registered trademarks of their respective owners.

* Responder rate = GLS = 0 or 1 at Maximum Frown Investigator Assessment

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