



ANTARES PHARMA ANNOUNCES MULTIPLE PRESENTATIONS AT THE SEXUAL MEDICINE SOCIETY SCIENTIFIC ANNUAL MEETING

QUICKSHOT® TESTOSTERONE DATA HAS BEEN SELECTED FOR BOTH ORAL AND POSTER PRESENTATIONS

EWING, NJ, October 28, 2016 - Antares Pharma, Inc. (NASDAQ: ATRS) today announced that data from the 52 week phase 3 study of the pharmacokinetics and safety of subcutaneous testosterone enanthate delivered through the QuickShot® auto injector was selected for an oral podium presentation at the 22nd Annual Fall Scientific Meeting of the Sexual Medicine Society of North America, which will be held at The Phoenician, Scottsdale, Arizona November 3rd through November 6th. A moderated poster presentation featuring additional data from the same 52 week phase 3 study which tracked psychosexual function in hypogonadal men will also be presented at the same meeting on November 3, 2016.

Oral Podium Presentation

The abstract, entitled “Safety and efficacy results from the phase 3, double blind, multicenter STEADY trial of a novel, pre-filled, subcutaneous (SC) auto injector for testosterone (T) replacement therapy,” was authored by Ronald S. Swerdloff, MD, Los Angeles Biomedical Research Institute and Harbor-UCLA Medical Center, Los Angeles, CA, et al.

The details for Dr. Swerdloff’s podium presentation are as follows:

Date: Thursday, November 3, 2016

Session: Oral Presentations: Hormones

Session Time: 4:30 p.m. MT

Location: The Phoenician, Scottsdale Arizona

Moderated Poster Presentation

The poster entitled “Improvements in psychosexual function among hypogonadal men enrolled in the STEADY trial of a novel, subcutaneous auto injector for testosterone replacement” was authored by Christina Wang, MD, Los Angeles Biomedical Research Institute and Harbor-UCLA Medical Center, Los Angeles, CA, et al.

The details for Dr. Wang’s moderated poster presentation are as follows:

Date: Thursday, November 3, 2016

Session: Moderated Poster Presentations: Hormones

Session Time: 5:30 p.m. MT

Location: The Phoenician, Scottsdale Arizona

About Antares Pharma

Antares Pharma focuses on self-administered parenteral pharmaceutical products. The Company’s product, OTREXUP™ (methotrexate) injection for subcutaneous use, is approved in the U.S. for the treatment of adults with severe active rheumatoid arthritis, children with active polyarticular juvenile idiopathic arthritis and adults with severe recalcitrant psoriasis. The Company and Teva

Pharmaceutical Industries, Ltd. (Teva) recently announced the commercial launch of VIBEX® Sumatriptan Injection USP for the acute treatment of migraine and cluster headache in the United States. Antares Pharma is also developing QuickShot® Testosterone for testosterone replacement therapy. The Company's technology platforms include VIBEX® disposable auto injectors, disposable multi-use pen injectors and reusable needle-free injectors. Antares Pharma has a multi-product deal with Teva that includes VIBEX® epinephrine, exenatide multi-dose pen, and teriparatide multi-dose pen. Our reusable needle-free injector for use with human growth hormone (hGH) is sold worldwide by Ferring B.V. The Company is also working with AMAG Pharmaceuticals on a subcutaneous method of administering Makena, a progesterone product indicated for use in lowering the risk of pre-term birth. For more information, [visit www.antaresspharma.com](http://www.antaresspharma.com).

SAFE HARBOR STATEMENT UNDER THE PRIVATE SECURITIES LITIGATION REFORM ACT OF 1995

This press release contains forward-looking statements within the meaning of the safe harbor provisions of the Private Securities Litigation Reform Act of 1995. Forward-looking statements are subject to certain risks and uncertainties that can cause actual results to differ materially from those described. Factors that may cause such differences include, but are not limited to: the timing and results of the phase 3 studies for QuickShot® Testosterone (QST) and acceptance of the data by the U.S. Food and Drug Administration ("FDA"); the timing and Company's ability to successfully complete a New Drug Application ("NDA") for QST, acceptance of the NDA for QST by the FDA and approval of the same by the FDA; Teva's ability to adequately and timely respond to the Complete Response Letter received from the FDA for the VIBEX® epinephrine pen ANDA and approval by the FDA of the same, the timing and therapeutic equivalence rating thereof, and any future purchase orders and revenue pre or post FDA approval; FDA action with respect to Teva's ANDA filed for the Exenatide pen and future revenue from the same; Teva's ability to successfully commercialize VIBEX® Sumatriptan Injection USP and the amount of revenue from the same; the outcome of the pending patent litigation between Teva Pharmaceutical Industries, Ltd. (Teva) and Eli Lilly and Company regarding the Teriparatide multi-dose pen; the timing and approval, if any, by the FDA of Teva's ANDA for the Teriparatide multi-dose pen and any future revenue resulting therefrom; continued growth of prescriptions and sales of OTREXUP™; the timing and results of the development project with AMAG Pharmaceuticals for an auto injector for Makena; the timing and results of research projects, clinical trials, and product candidates in development; actions by the FDA or other regulatory agencies with the respect to the Company's products or product candidates of its partners; continued growth in product, development, licensing and royalty revenue; the Company's ability to obtain financial and other resources for its research, development, clinical, and commercial activities and other statements regarding matters that are not historical facts, and involve predictions. These statements involve known and unknown risks, uncertainties and other factors that may cause actual results, performance, achievements or prospects to be materially different from any future results, performance, achievements or prospects expressed in or implied by such forward-looking statements. In some cases you can identify forward-looking statements by terminology such as "may", "will", "should", "would", "expect", "intend", "plan", "anticipate", "believe", "estimate", "predict", "potential", "seem", "seek", "future", "continue", or "appear" or the negative of these terms or similar expressions, although not all forward-looking statements contain these identifying words. Additional information concerning these and other factors that may cause actual results to differ materially from those anticipated in the forward-looking statements is contained in the "Risk Factors" section of the Company's Annual Report on Form 10-K for the year ended December 31, 2015, and in the Company's other periodic reports and filings with the Securities and Exchange Commission. The Company cautions investors not to place undue reliance on the forward-looking statements contained in this press release. All forward-looking statements are based on information currently available to the Company on the date hereof, and the Company undertakes no obligation to revise or update these forward-looking statements to reflect events or circumstances after the date of this press release, except as required by law.

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