NEWS RELEASE

ANTARES PHARMA PROVIDES REGULATORY UPDATE ON SUMATRIPTAN INJECTION USP

EWING, NJ, January 26, 2015 -- Antares Pharma, Inc. (NASDAQ: ATRS) today announced that the U.S. Food and Drug Administration (FDA) has issued a complete response letter regarding the Abbreviated New Drug Application (ANDA) for Sumatriptan Injection USP for the acute treatment of migraine. The complete response letter from the FDA provided revisions to labelling and cited minor deficiencies. If approved, Sumatriptan Injection USP would represent the Company's first ANDA approval and second device approved from the VIBEX® auto injector platform. As previously disclosed, Teva Pharmaceutical Industries, Ltd. would distribute the product and share the profits equally with Antares Pharma.

Eamonn P. Hobbs, President and Chief Executive Officer, stated, "We are pleased with the feedback received from the FDA in the complete response letter. The FDA has outlined the steps necessary to support approval of the ANDA, and we plan to work closely with the Agency on a response."

About Antares Pharma

Antares Pharma focuses on self-administered parenteral pharmaceutical products. The Company markets OTREXUP™ (methotrexate) injection for subcutaneous use in the treatment of adults with severe active rheumatoid arthritis and children with active polyarticular juvenile idiopathic arthritis. LEO Pharma markets OTREXUP™ to dermatologists for adults with severe recalcitrant psoriasis. Antares Pharma is also developing QuickShot® Testosterone for testosterone replacement therapy, and VIBEX® Sumatriptan for the acute treatment of migraines. 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 another undisclosed multi-dose pen. Our reusable needle-free injector for use with human growth hormone (hGH) is sold worldwide by Ferring Pharmaceuticals BV.

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, including statements made with respect to the Company's Abbreviated New Drug Application for Sumatriptan Injection USP, the approval thereof and Teva's distribution thereof; 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. Such forward-looking statements are not guarantees of future performance and are subject to risks and uncertainties that may cause actual results to differ materially from those anticipated by the forward-looking statements. 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, 2013, 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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