Mylan Receives FDA Approval for Mirtazapine Tablets

PITTSBURGH, Jun 19, 2003 (BUSINESS WIRE) -- Mylan Laboratories Inc. (NYSE:MYL) announced today that the U.S. Food and Drug Administration has granted final approval for its Abbreviated New Drug Application for Mirtazapine Tablets in 15 mg, 30 mg and 45 mg strengths. Mirtazapine is the generic version of Organon Inc.'s, Remeron®, which is indicated for the treatment of major depressive disorder.

Mylan will be launching Mirtazapine immediately.

Mylan Laboratories Inc. is a leading pharmaceutical company that develops, manufactures and markets generic and proprietary prescription products. Mylan has two operating segments that market an extensive line of generic and branded products through four business units: Mylan Pharmaceuticals Inc., Mylan Technologies Inc., UDL Laboratories, Inc. and Bertek Pharmaceuticals Inc. For more information about Mylan, visit www.mylan.com.

This press release may contain forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995. Because such statements are subject to risks and uncertainties, actual future results may differ materially from those expressed or implied by such forward-looking statements. We refer you to the risk factors and other disclosures contained in our periodic SEC filings. We undertake no duty to update our forward-looking statements, even though our situation may change in the future.

SOURCE: Mylan Laboratories Inc.

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