Mylan Receives FDA Approval for Midodrine Hydrochloride Tablets

PITTSBURGH, Sep 11, 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 Midodrine Hydrochloride Tablets in 2.5 mg, 5 mg and 10 mg strengths. Midodrine Hydrochloride is the generic version of Shire Pharmaceuticals ProAmatine®.

Mylan will be launching Midodrine Hydrochloride Tablets immediately.

Mylan Laboratories Inc. is a leading pharmaceutical company with four subsidiaries, Mylan Pharmaceuticals Inc., Mylan Technologies Inc., UDL Laboratories Inc. and Bertek Pharmaceuticals Inc. that develop, manufacture and market an extensive line of generic and proprietary products.

For more information about Mylan, visit www.mylan.com.

This press release may contain forward-looking statements, including with regard to the sale of Midodrine Hydrochloride Tablets. These statements are made pursuant to the safe harbor provisions of the Private Securities Litigation Reform Act of 1995. Because such statements inherently involve risks and uncertainties, actual future results may differ materially from those expressed or implied by such forward-looking statements. Factors that could cause or contribute to such differences include, but are not limited to: the possible negative effects of any interruption of manufacture of the Tablets at the Company's facility; uncertainties regarding market acceptance and demand for Midodrine Hydrochloride; dependence on third-party suppliers and distributors for raw materials; and the other risks detailed in the Company's filings with the Securities and Exchange Commission. The Company undertakes no obligation to update these statements, for revisions or changes after the date of this release.

SOURCE: Mylan Laboratories Inc.

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