# Mylan Announces Final FDA Approval for Cetirizine Hydrochloride Tablets

PITTSBURGH, Dec 28, 2007 /PRNewswire-FirstCall via COMTEX News Network/ -- Mylan Inc. (NYSE: MYL) today announced that Mylan Pharmaceuticals Inc. has received final approval from the U.S. Food and Drug Administration (FDA) for its Abbreviated New Drug Application (ANDA) for Cetirizine Hydrochloride Tablets (OTC), 5 mg and 10 mg.

Cetirizine HCl Tablets are the generic version of Pfizer's Zyrtec® Tablets, which had U.S. sales of approximately $1.4 billion for the 12 months ending Sept. 30, 2007.

This product will be shipped immediately.

Mylan Inc. is one of the world's leading quality generic and specialty pharmaceutical companies. The Company offers one of the industry's broadest and highest quality product portfolios, a robust product pipeline and a global commercial footprint through operations in more than 90 countries. Through its controlling interest in Matrix Laboratories Limited, Mylan has direct access to one of the largest active pharmaceutical ingredient (API) manufacturers in the world. Dey L.P., Mylan's fully integrated specialty business, provides the Company with innovative and diversified opportunities in the respiratory and allergy therapeutic areas.

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