Mylan Inc. Announces Tentative FDA Approval for Lamotrigine Tablets

PITTSBURGH, Dec. 12 /PRNewswire-FirstCall/ -- Mylan Inc. (NYSE: MYL) today announced that Mylan Pharmaceuticals Inc. has received tentative approval from the U.S. Food and Drug Administration (FDA) for its Abbreviated New Drug Application (ANDA) for Lamotrigine Tablets, 25 mg, 100 mg, 150 mg and 200 mg.

Lamotrigine Tablets are the generic version of GlaxoSmithKline's Lamictal® Tablets, which had U.S. sales of approximately $1.97 million for the 12 months ending Sept. 30, 2007.

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.

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

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