Mylan Receives Approval for Generic Version of Wellbutrin SR(R)

PITTSBURGH, April 12, 2010 /PRNewswire via COMTEX News Network/ -- Mylan Inc. (Nasdaq: MYL) today announced that its subsidiary Mylan Pharmaceuticals Inc. received final approval from the U.S. Food and Drug Administration (FDA) for its Abbreviated New Drug Application (ANDA) for Bupropion Hydrochloride Extended-Release Tablets USP, (SR), 100 mg, 150 mg and 200 mg, the generic version of GlaxoSmithKline's antidepressant Wellbutrin SR(R).

Bupropion Hydrochloride Extended-Release Tablets had U.S. sales of approximately $363 million for the 12 months ending Dec. 31, 2009, according to IMS Health. Mylan has started shipping this product.

Currently, Mylan has 141 ANDAs pending FDA approval representing $96.2 billion in annual brand sales, according to IMS Health. Thirty-eight of these pending ANDAs are potential first-to-file opportunities, representing $19.8 billion in annual brand sales, for the 12 months ending Dec. 31, 2009, according to IMS Health.

Mylan Inc. ranks among the leading generic and specialty pharmaceutical companies in the world and provides products to customers in more than 140 countries and territories. The company maintains one of the industry's broadest and highest quality product portfolios supported by a robust product pipeline; operates one of the world's largest active pharmaceutical ingredient manufacturers; and runs a specialty business focused on respiratory, allergy and psychiatric therapies. For more information, please visit www.mylan.com.

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