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Mylan Launches Cabergoline Tablets

PITTSBURGH, Dec. 5, 2013 /PRNewswire/ -- Mylan Inc. (Nasdaq: MYL) today announced that its U.S.-based subsidiary Mylan Pharmaceuticals Inc. has launched Cabergoline Tablets USP, 0.5 mg. Cabergoline Tablets USP are indicated for the treatment of hyperprolactinemic disorders, either idiopathic or due to pituitary adenomas. Mylan received final approval from the U.S. Food and Drug Administration (FDA) for its Abbreviated New Drug Application (ANDA) for this product.

Cabergoline Tablets USP, 0.5 mg, had U.S. sales of approximately $41.67 million for the 12 months ending Sept. 30, 2013, according to IMS Health.

Currently, Mylan has 178 ANDAs pending FDA approval representing $90.3 billion in annual sales, according to IMS Health. Forty of these pending ANDAs are potential first-to-file opportunities, representing $24.1 billion in annual brand sales, for the 12 months ending June 30, 2013, according to IMS Health.

Mylan is a global pharmaceutical company committed to setting new standards in health care. Working together around the world to provide 7 billion people access to high quality medicine, we innovate to satisfy unmet needs; make reliability and service excellence a habit; do what's right, not what's easy; and impact the future through passionate global leadership. We offer a growing portfolio of more than 1,200 generic pharmaceuticals and several brand medications. In addition, we offer a wide range of antiretroviral therapies, upon which approximately 40% of HIV/AIDS patients in developing countries depend. We also operate one of the largest active pharmaceutical ingredient manufacturers and currently market products in approximately 140 countries and territories. Our workforce of more than 20,000 people is dedicated to improving the customer experience and increasing pharmaceutical access to consumers around the world. But don't take our word for it. See for yourself. See inside. mylan.com

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