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Mylan Launches Thiamine Hydrochloride Injection

PITTSBURGH, June 28, 2012 /PRNewswire/ -- Mylan Inc. (Nasdaq: MYL) today announced that its Mylan Institutional business has received final approval from the U.S. Food and Drug Administration (FDA) for its Abbreviated New Drug Application (ANDA) for preservative-free Thiamine Hydrochloride Injection, 100 mg/mL, packaged in 200 mg/2 mL multiple-dose vials. This product is indicated for the treatment of thiamine deficiency.

Thiamine Hydrochloride Injection, 100 mg/mL, had U.S. sales of approximately $18.6 million for the 12 months ending March 31, 2012, according to IMS Health. Mylan is shipping this product, presented in 25-vial packs, immediately.

Currently, Mylan has 169 ANDAs pending FDA approval representing $83.9 billion in annual sales, according to IMS Health. Thirty-seven of these pending ANDAs are potential first-to-file opportunities, representing $25.6 billion in annual brand sales, for the 12 months ending Dec. 31, 2011, 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 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 1,100 generic pharmaceuticals and several brand medications. In addition, we offer a wide range of antiretroviral therapies, upon which approximately one-third 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 150 countries and territories. Our workforce of more than 18,000 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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