Mylan Announces Final FDA Approval for Zaleplon Capsules

PITTSBURGH, June 9 /PRNewswire-FirstCall/ -- Mylan Inc. (NYSE: MYL) today announced that its subsidiary, Genpharm ULC, has received final approval from the U.S. Food and Drug Administration (FDA) for its Abbreviated New Drug Application (ANDA) for Zaleplon Capsules, 5 mg and 10 mg.

Zaleplon Capsules are the generic version of King Pharmaceuticals' Sonata® Capsules, which had U.S. sales of approximately $88 million for the 12 months ending March 31, 2008, according to IMS Health.

This product is shipping immediately and will be sold under the Mylan Pharmaceuticals brand.

Mylan Inc., with a presence in more than 90 countries, ranks among the leading diversified generic and specialty pharmaceutical companies in the world. The company maintains one of the industry's broadest -- and highest quality -- product portfolios, supported by a robust product pipeline; owns a controlling interest in the world's second largest active pharmaceutical ingredient manufacturer; and operates a specialty business focused on respiratory and allergy therapies.

SOURCE Mylan Inc. 06/09/2008 CONTACT: Media, Michael Laffin, or Investors, Kris King, both of Mylan Inc., +1-724-514-1813 Web site: http://www.mylan.com (MYL)