FOR IMMEDIATE RELEASE

AMNEAL RECEIVES FDA APPROVAL FOR PRIMIDONE TABLETS, USP

Paterson, New Jersey (USA), March 10, 2008 – Amneal Pharmaceuticals is pleased to announce that it received US FDA approval to manufacture Primidone Tablets, USP in strengths of 50mg and 250mg, effective March 1, 2008. Primidone is an AB-Rated, therapeutically equivalent alternative to Mysoline® (a trademark of Valeant Pharmaceuticals).

The Primidone approval is Amneal's 17th ANDA approval and continues to build the momentum for the aggressive growth Amneal is experiencing. Separately, in December, Amneal closed on the acquisition of a liquid Rx manufacturing plant in New Jersey and is currently negotiating to acquire several approved ANDAs.

Primidone, used alone or concomitantly with other anticonvulsants, is indicated in the control of grand mal, psychomotor, and focal epileptic seizures. It may control grand mal seizures refractory to other anticonvulsant therapy. Amneal will begin shipping Primidone Tablets in the 50 and 250mg strengths as of April 1, 2008 and will have the product available through wholesalers-distributors as well as directly to the trade.

Amneal Pharmaceuticals LLC, headquartered in Paterson, NJ, is a USA-based firm that develops, manufacturers and distributes generic pharmaceutical products regulated and approved by the US FDA. Positioned as "Generic's New Generation," the company utilizes diverse R&D and manufacturing expertise to conceive breakthrough developments with lasting impact. Vigorous ANDA growth and broad product acquisitions are key features of Amneal's strategic growth plan, as is the company's commitment to building deep relationships with its customer base. Amneal delivers superior service levels, quality products, and dynamic value throughout the pharmaceutical industry.

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Brand Name:

Orange Book Rating: AB rated

Strengths: 50mg and 250mg

Indication: