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FOR IMMEDIATE RELEASE

Sun Pharma announces USFDA approval for generic Razadyne® ER

Mumbai, February 03, 2011: Sun Pharmaceutical Industries Ltd. (Reuters: SUN.BO, Bloomberg: SUNP IN, NSE:SUNPHARMA, BSE: 524715) announced that USFDA has granted its subsidiary an approval for an Abbreviated New Drug Application (ANDA) to market a generic version of Razadyne® ER, galantamine hydrobromide extended-release capsules.

These generic galantamine hydrobromide extended-release capsules, 8 mg (base), 16 mg (base) and 24 mg (base) are indicated in the treatment of mild to moderate dementia of the Alzheimer's type.

Razadyne® ER has annual sale of approximately \$ 50 million in the US.

Razadyne® is a registered trademark of Ortho-McNeil Janssen Pharmaceuticals, Inc.

About Sun Pharmaceutical Industries Ltd.

Established in 1983, listed since 1994 and headquartered in India, Sun Pharmaceutical Industries Ltd. (Reuters: SUN.BO, Bloomberg: SUNP IN, NSE: SUNPHARMA, BSE: 524715) is an international, integrated, speciality pharmaceutical company. It manufactures and markets a large basket of pharmaceutical formulations as branded generics as well as generics in India, US and several other markets across the world. In India, the company is a leader in niche therapy areas of psychiatry, neurology, cardiology, diabetology, gastroenterology, ophthalmology and orthopedics. The company has strong skills in product development, process chemistry, and manufacturing of complex API, as well as dosage forms. More information about the company can be found at www.sunpharma.com.

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