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FOR IMMEDIATE RELEASE

Sun Pharma announces USFDA approval for generic Zyprexa® tablets

Mumbai, April 24, 2012: Sun Pharmaceutical Industries Ltd. (Reuters: SUN.BO, Bloomberg: SUNP IN, NSE: SUNPHARMA, BSE: 524715) announced that USFDA has granted an approval for its Abbreviated New Drug Application (ANDA) for generic version of Zyprexa®, Olanzapine Tablets USP, 2.5 mg, 5 mg, 7.5 mg, 10 mg, 15 mg and 20 mg.

These generic Olanzapine tablets, 2.5 mg, 5 mg, 7.5 mg, 10 mg, 15 mg and 20 mg are therapeutic equivalents of Eli Lilly's Zyprexa® tablets. Olanzapine tablets have annual sale of approximately $ 3.3 billion in the US.

Olanzapine tablets are indicated for the treatment of schizophrenia, bipolar I disorder (manic or mixed episodes).

Zyprexa® is a registered trademark of Eli Lilly & Co.

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, nephrology, 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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