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FOR IMMEDIATE RELEASE

Sun Pharma announces US FDA approval for generic Cymbalta®

Mumbai, December 12, 2013: Sun Pharmaceutical Industries Ltd. (Reuters: SUN.BO, Bloomberg: SUNP IN, NSE: SUNPHARMA, BSE: 524715) today announced that the US FDA has granted its subsidiary final approval for its Abbreviated New Drug Application (ANDA) to market a generic version of Cymbalta®, Duloxetine Delayed-Release Capsules USP, 20 mg, 30 mg and 60 mg.

Duloxetine Delayed-Release Capsules USP, 20 mg, 30 mg and 60 mg are therapeutic equivalents of Eli Lilly & Company's Cymbalta® Delayed-Release Capsules. These Capsules have annual sales of approximately USD 5.5 billion in the US. Duloxetine Delayed-Release Capsules USP are indicated for the treatment of major depressive disorder (MDD), generalized anxiety disorder (GAD) and diabetic peripheral neuropathic pain (DPNP).

Sun Pharma's subsidiary, being one of the first-to-file ANDAs for generic Cymbalta® with a para IV certification, is eligible for shared 180-day marketing exclusivity in the US.

Cymbalta® is a registered trademark of Eli Lilly & Company, Inc.

About Sun Pharma

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 specialty pharmaceutical company with over 70% sales from global markets. It manufactures and markets a large basket of pharmaceutical formulations as branded generics as well as generics in US, India and several other markets across the world. For the year ending March 2013, overall revenues were at US$2.1 billion, of which US contributed US$1.1 billion. In India, the company is a leader in niche therapy areas of psychiatry, neurology, cardiology, nephrology, gastroenterology, orthopedics and ophthalmology. The company has strong skills in product development, process chemistry, and manufacturing of complex dosage forms. More information about the company can be found at www.sunpharma.com.

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