17/B, Mahal Industrial Estate,
Mahakali Caves Road,
Andheri (East), Mumbai 400 093 India
Tel.: (91-22) 6645 5645
Fax.: (91-22) 6645 5685

FOR IMMEDIATE RELEASE

Sun Pharma announces USFDA tentative approval for generic Cymbalta®

Mumbai, November 24, 2010: Sun Pharmaceutical Industries Ltd. (Reuters: SUN.BO, Bloomberg: SUNP IN, NSE:SUNPHARMA, BSE: 524715) announced that USFDA has granted its subsidiary a tentative approval for an Abbreviated New Drug Application (ANDA) to market a generic version of Cymbalta®, duloxetine hydrochloride delayed-release capsules.

These generic duloxetine hydrochloride delayed-release capsules, 20 mg (base), 30 mg (base), and 60 mg (base) are indicated in the treatment of Major Depressive Disorder (MDD), Generalized Anxiety Disorder (GAD) and Diabetic Peripheral Neuropathic Pain (DPNP)

Cymbalta® has annual sale of approximately $ 3 billion in the US.

Cymbalta® is a registered trademark of Eli Lilly & Company

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, orthopedics and ophthalmology. 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.

Contacts

Uday Baldota
Tel +91 22 6645 5645, Xtn 605
Tel Direct +91 22 66455605
Mobile +91 98670 10529
E mail uday.baldota@sunpharma.com

Mira Desai
Tel +91 22 6645 5645, Xtn 606
Tel Direct +91 22 66455606
Mobile +91 98219 23797
E mail mira.desai@sunpharma.com