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FOR IMMEDIATE RELEASE

Sun Pharma announces US FDA approval for generic Temodar®

Mumbai, February 13, 2014: 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 Temodar®, Temozolomide Capsules, 5 mg, 20 mg, 100 mg, 140 mg, 180 mg and 250 mg.

Temozolomide Capsules, 5 mg, 20 mg, 100 mg, 140 mg, 180 mg and 250 mg are therapeutic equivalents of Merck Sharp & Dohme Corporation's Temodar® Capsules. These Capsules have annual sales of approximately USD 400 million in the US.

Temozolomide Capsules are indicated for the treatment of adult patients with newly diagnosed glioblastoma multiforme concomitantly with radiotherapy and then as maintenance treatment and for adult patients with refractory anaplastic astrocytoma who have experienced disease progression on a drug regimen containing nitrosourea and procarbazine.

Temodar® is the registered trademark of Merck Sharp & Dohme Corp.

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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