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FOR IMMEDIATE RELEASE

Sun Pharma announces USFDA approval for generic Doxil®

Mumbai, February 05, 2013: Sun Pharmaceutical Industries Ltd. (Reuters: SUN.BO, Bloomberg: SUNP IN, NSE: SUNPHARMA, BSE: 524715) today announced that USFDA has granted its subsidiary an approval for its Abbreviated New Drug Application (ANDA) for generic version of Doxil®, Doxorubicin HCl Liposome Injection USP, 2mg/mL, packaged in 20 mg/10mL and 50 mg/25mL single-use vials.

This generic Doxorubicin HCl Liposome injection USP, 2 mg/mL is therapeutically equivalent to Doxil® Liposome Injection, 2 mg/mL of Janssen Research and Development, LLC.

Doxorubicin HCl liposome injection is used to treat patients with ovarian cancer that has progressed or recurred after platinum-based chemotherapy.

DOXIL® is a registered trademark of Janssen Products, LP (USA).

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, 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.

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