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FOR IMMEDIATE RELEASE

Sun Pharma announces USFDA approval for generic Accupril® tablets

Mumbai, June 19, 2009: 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 Accupril®, quinapril hydrochloride tablets.

These quinapril hydrochloride tablets, therapeutically equivalent to Accupril® tablets from Pfizer, are available in four strengths viz. 5 mg, 10 mg, 20 mg and 40 mg and have annual sales of approximately USD 45 million in the US.

Quinapril is an antihypertensive.

Accupril® is a registered trademark of Pfizer.

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, diabetology, gastroenterology, 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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