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FOR IMMEDIATE RELEASE

Sun Pharma announces USFDA tentative approval for generic Uroxatral Extended Release ® tablets

Mumbai, May 7, 2009: 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 its Abbreviated New Drug Application (ANDA) for generic Uroxatral Extended Release ® tablets.

These alfuzosin hydrochloride 10 mg tablets are therapeutically equivalent to Uroxatral Extended Release® tablets from sanofi-aventis. Alfuzosin hydrochloride tablets have annual sales of approximately USD 180 million in the US.

Alfuzosin is an alpha 1 blocker for the treatment of the signs and symptoms of BPH.

Uroxatral® is a registered trademark of sanofi aventis U.S. LLC

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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