

FOR IMMEDIATE RELEASE

Sun Pharma Announces US FDA Approval for Generic Amphotericin B Liposome Injection

Mumbai, India, December 15, 2021: Sun Pharmaceutical Industries Limited (Reuters: SUN.BO, Bloomberg: SUNP IN, NSE: SUNPHARMA, BSE: 524715, "Sun Pharma" including its subsidiaries and/or associate companies), today announced that one of its wholly-owned subsidiaries has received final approval from US FDA for its Abbreviated New Drug Application (ANDA) for generic Amphotericin B Liposome for Injection, 50 mg/vial Single-Dose Vial. The generic product approval is based on AmBisome® Liposome for Injection, 50 mg/vial as a reference product.

Sun Pharma has been granted Competitive Generic Therapy (CGT) designation by US FDA and being the first approved generic, is eligible for 180 days of CGT exclusivity for the product.

As per October 2021 IQVIA Health data, AmBisome® Liposome for Injection, 50mg/vial had annualized sales of approximately US\$ 136 million in USA.

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About Sun Pharmaceutical Industries Limited (CIN - L24230GJ1993PLC019050):

Sun Pharma is the world's fourth largest specialty generic pharmaceutical company and India's top pharmaceutical company. A vertically integrated business and a skilled team enables it to deliver high-quality products, trusted by customers and patients in over 100 countries across the world, at affordable prices. Its global presence is supported by manufacturing facilities spread across 6 continents and approved by multiple regulatory agencies, coupled with a multi-cultural workforce comprising over 50 nationalities. Sun Pharma fosters excellence through innovation supported by strong R&D capabilities across multiple R&D centers, with investments of approximately 6-7% of annual revenues in R&D. For further information, please visit www.sunpharma.com & follow us on [Twitter@SunPharma_Live](https://twitter.com/SunPharma_Live).

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