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Dr. Reddy's Laboratories announces the launch of Fosaprepitant for Injection in the U.S. Market

Hyderabad, India, September 11, 2019

For Immediate Release

Hyderabad, India and Princeton, NJ, USA. September 11, 2019— Dr. Reddy's Laboratories Ltd. (BSE: 500124, NSE: DRREDDY, NYSE: RDY, along with its subsidiaries together referred to as "Dr. Reddy's") today announced the launch of Fosaprepitant for Injection, the therapeutic generic equivalent of EMEND® (fosaprepitant) for injection, approved by the U.S. Food and Drug Administration (USFDA).

"We are pleased to be among the first generics to launch this product," explains Marc Kikuchi, Chief Executive Officer, North America Generics, Dr. Reddy's Laboratories. "This is a great addition to our injectable portfolio in the U.S. market as we continue to augment our product offering and drive growth for the hospital segment."

The EMEND® for Injection brand had an U.S. sales of approximately \$279 million MAT for the most recent twelve months ending in July 2019 according to IQVIA Health*.

Dr. Reddy's Fosaprepitant for Injection is available in 150 mg single-dose vial for reconstitution.

EMEND® is a trademark of Merck Sharp & Dohme Corp.

*IQVIA Retail and Non-Retail MAT July 2019

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About Dr. Reddy's: Dr. Reddy's Laboratories Ltd. (BSE: 500124, NSE: DRREDDY, NYSE: RDY) is an integrated pharmaceutical company, committed to providing affordable and innovative medicines for healthier lives. Through its three businesses - Pharmaceutical Services & Active Ingredients, Global Generics and Proprietary Products – Dr. Reddy's offers a portfolio of products and services including APIs, custom pharmaceutical services, generics, biosimilars and differentiated formulations. Our major therapeutic areas of focus are gastrointestinal, cardiovascular, diabetology, oncology, pain management and dermatology. Dr. Reddy's operates in markets across the globe. Our major markets include – USA, India, Russia & CIS countries, and Europe. For more information, log on to: www.drreddys.com

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