Press Release **For Immediate Release**

Glenmark Generics receives final ANDA approval for Omeprazole DR Capsules

Mumbai – India, November 5, 2014: Glenmark Generics Inc., USA the subsidiary of Glenmark Generics Limited has been granted final approval for its abbreviated new drug application (ANDA) from the United States Food and Drug Administration (U.S. FDA) for Omeprazole Delayed Release Capsules, their generic version of Prilosec® by AstraZeneca.

Omeprazole DR Capsules are indicated for the short-term treatment of active duodenal ulcer in adults. The approval is for the 10, 20, and 40mg strengths of Omeprazole. According to IMS Health sales data for the 12 month period ending September 2014, Omeprazole garnered annual sales of approximately USD 520 million.

Glenmark's current portfolio consists of 94 products authorized for distribution in the U.S. marketplace and 72 ANDA's pending approval with the U.S. FDA. In addition to these internal filings, GGI continues to identify and explore external development partnerships to supplement and accelerate the growth of the existing pipeline and portfolio.

About Glenmark Generics Limited:

Glenmark Generics Limited (GGL) is a subsidiary of Glenmark Pharmaceuticals Limited (Glenmark) and aims to be a global integrated Generic and API leader. GGL has an established presence in North America and developing an EU presence. It primarily sells its FDF products in the United States ("US") and the European Union ("EU"), as well as its oncology FDF products in South America. The Company supplies APIs to customers in approximately 80 countries, including the US, various countries in the EU, South America and India.

For further information, please contact:

Jason D'Souza / Rajdeep Barooah
Glenmark, Mumbai, India Tel: [+91 22] 40189919/984
Email: corpcomm@glenmarkpharma.com