

Press Release

For Immediate Release

Glenmark Generics receives final ANDA approval for Eszopiclone Tablets

April 16, 2014: Glenmark Generics Inc., USA a subsidiary of Glenmark Generics Limited has been granted final abbreviated new drug approval (ANDA) from the United States Food and Drug Administration (US FDA) for Eszopiclone Tablets. Glenmark will commence distribution of the product immediately.

Eszopiclone Tablets are Glenmark's generic version of Sunovion's Lunesta®. Eszopiclone is indicated for the treatment of insomnia. The approval is for the 1mg, 2mg and 3mg tablets. For the 12 month period ending December 2013, Lunesta® garnered annual sales of USD 824 Million according to IMS Health.

Glenmark's current portfolio consists of 91 products authorized for distribution in the U.S. marketplace and 63 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.

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