

Glenmark receives ANDA approval for Clobetasol Propionate Foam

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It is a generic version of Olux[®]1 Foam, 0.05%, of Mylan Pharmaceuticals



Glenmark Pharmaceuticals has been granted final approval by the United States Food & Drug Administration (U.S. FDA) for Clobetasol Propionate Foam, 0.05%, a generic version of Olux[®]1 Foam, 0.05%, of Mylan Pharmaceuticals, Inc.

According to IQVIATM sales data for the 12 month period ending December 2018, the Olux[®] Foam, 0.05% market² achieved annual sales of approximately \$50.9 million.

Glenmark's current portfolio consists of 150 products authorized for distribution in the U.S. marketplace and 52 ANDA's pending approval with the U.S. FDA.