

LUPIN GETS FDA'S NOD FOR DRUG MARKETING

New Delhi: Lupin has received tentative approval from the US health regulator to market a generic version of Prezista tablets, used in treating HIV infection. The company has received tentative approval from the United States Food and Drugs Administration (FDA) to market its Darunavir

Ethanolate tablets in multiple strengths in the American market, Lupin said in a statement. Darunavir tablets are indicated for the treatment of HIV-

1 infections. According to IMS MAT September, 2014 sales data, Janssen's Prezista tablets had annual US sales of \$1.2 billion. Last week, the Mumbai-based firm had received US health regulator's approval to market a generic version of ViiV Healthcare's Epivir Tablets, used in treating HIV infection, in the American market. Lupin shares closed at ₹1,428.70 apiece on the BSE, down 0.21 per cent from its previous close. — PTI



4
11/1/2015
BSE

Dir (Sec)

Handwritten signature
21/1/15

Comptroller (Gen)

TO, NIC

Handwritten signature
6.1.15

Company