

January 10, 2023

BSE Limited Code: 532321

1st Floor, P J Towers, Dalal Street, <u>Mumbai-400001</u>

National Stock Exchange of India Limited

Exchange Plaza, 5th Floor, Plot No. C/1, G Block, Bandra-Kurla Complex, Bandra (East), <u>Mumbai-400051</u>

Re.: Press Release

Dear Sir / Madam,

Please find enclosed a copy of press release dated January 10, 2023 titled "Zydus receives final approval from the USFDA for Febuxostat Tablets".

The contents of the press release give full details.

Please bring the aforesaid news to the notice of the members of the exchange and the investors' at large.

Thanking you,

Yours faithfully,
For, ZYDUS LIFESCIENCES LIMITED

DHAVAL N. SONI COMPANY SECRETARY

Encl.: As above



Code: Zyduslife



Zydus receives final approval from the USFDA for Febuxostat Tablets

Ahmedabad, India, 10 January, 2023

Zydus Lifesciences Limited (formerly known as Cadila Healthcare Limited) has received final approval from the United States Food and Drug Administration (USFDA) to market Febuxostat Tablets, 40 mg and 80 mg (USRLD: Uloric Tablets).

Febuxostat tablets are indicated to lower hyperuricemia (high uric acid in the blood) in patients with gout who have been treated with allopurinol that did not work well or cannot be treated with allopurinol. The drug will be manufactured at the group's formulation manufacturing facility at Moraiya, Ahmedabad, India.

Febuxostat tablets had annual sales of USD 32 million in the United States according to IQVIA data (IQVIA MAT Sept. 2022).

The group now has 340 approvals and has so far filed over 431* ANDAs since the commencement of the filing process in FY 2003-04.

(*as of 30th September 2022)



For further information please contact: The Corporate Communications Department

Zydus Lifesciences Limited

(formerly known as Cadila Healthcare Limited)
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