

December 30, 2022

BSE Limited Code: 532321

1st Floor, P J Towers, Dalal Street, Mumbai-400001

National Stock Exchange of India Limited

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

Re.: Press Release

Dear Sir / Madam,

Please find enclosed a copy of press release dated December 30, 2022 titled "Zydus receives final approval from the USFDA for Triazolam Tablets".

Code: Zyduslife

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



Zydus receives final approval from the USFDA for Triazolam Tablets

Ahmedabad, India, 30 December, 2022

Zydus Lifesciences Limited (formerly known as Cadila Healthcare Limited) has received final approval from the United States Food and Drug Administration (USFDA) to market Triazolam Tablets USP, 0.125 mg and 0.25 mg (USRLD: HALCION® tablets).

Triazolam tablets are used on a short-term basis to treat insomnia (difficulty falling asleep or staying asleep). It works by slowing activity in the brain to allow sleep. The drug will be manufactured at the group's formulation manufacturing facility at Moraiya, Ahmedabad, India.

Triazolam tablets had annual sales of USD 11.7 million in the United States according to IQVIA data (IQVIA MAT Sept. 2022).

The group now has 339 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

CIN: L24230GJ1995PLC025878

(formerly known as Cadila Healthcare Limited)
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