

April 6, 2023

BSE Limited Code: 532321

P J Towers, Dalal Street, Mumbai-400001

National Stock Exchange of India Limited

Exchange Plaza, C/1, Block G, Bandra-Kurla Complex, Bandra (East), Mumbai-400051

Re.: Press Release

Dear Sir / Madam,

Please find enclosed copy of two press releases, both dated April 6, 2023, titled (1) "Zydus receives final approval from the USFDA for Acetazolamide Tablets USP, 125 mg and 250 mg" and (2) "Zydus receives final approval from the USFDA for Carbidopa and Levodopa Tablets USP, 10 mg/100 mg, 25 mg/100 mg, and 25 mg/250 mg".

The contents of the press releases 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 Acetazolamide Tablets USP, 125 mg and 250 mg

Ahmedabad, India, 6 April, 2023

Zydus Lifesciences Limited (including its subsidiaries/affiliates, hereafter referred to as "Zydus") has received final approval from the United States Food and Drug Administration (USFDA) to manufacture and market Acetazolamide Tablets USP, 125 mg and 250 mg (USRLD: Diamox Tablets, 125 mg and 250 mg).

Acetazolamide is used to treat glaucoma, a condition in which increased pressure in the eye can lead to gradual loss of vision. Acetazolamide decreases the pressure in the eye. Acetazolamide is also used to reduce the severity and duration of symptoms (upset stomach, headache, shortness of breath, dizziness, drowsiness, and fatigue) of altitude (mountain) sickness. The drug is used with other medicines to reduce edema (excess fluid retention) and to help control seizures in certain types of epilepsy. The product will be manufactured at the group's formulation manufacturing facility in Baddi, Himachal Pradesh (India).

Acetazolamide Tablets USP, 125 mg and 250 mg had annual sales of USD 16 mn in the United States (IQVIA MAT Dec. 2022).

The group now has 358 approvals and has so far filed over 440* ANDAs since the commencement of the filing process in FY 2003-04.

(*as of 31st December 2022)



For further information please contact: The Corporate Communications Department

Zydus Lifesciences Limited

(formerly known as Cadila Healthcare Limited)

Regd. Office : 'Zydus Corporate Park',

Scheme No. 63, Survey No. 536, Khoraj (Gandhinagar), Nr. Vaishnodevi Circle, S. G. Highway, Ahmedabad 382 481, Gujarat, India. | Phone: +91-79-71800000, +91-79-48040000 | website: www.zyduslife.com CIN: L24230GJ1995PLC025878



Zydus receives final approval from the USFDA for Carbidopa and Levodopa Tablets USP, 10 mg/100 mg, 25 mg/100 mg, and 25 mg/250 mg

Ahmedabad, India, 6 April, 2023

Zydus Lifesciences Limited (including its subsidiaries/affiliates, hereafter referred to as "Zydus") has received final approval from the United States Food and Drug Administration (USFDA) to manufacture and market Carbidopa and Levodopa Tablets USP, 10 mg/100 mg, 25 mg/100 mg, and 25 mg/250 mg (USRLD: Sinemet Tablets, 10 mg/100 mg, 25 mg/100 mg, and 25 mg/250 mg).

Carbidopa and Levodopa is used to treat symptoms of Parkinson's disease or Parkinson-like symptoms (such as shakiness, stiffness, difficulty moving). This medicine is also used to treat Parkinson symptoms caused by carbon monoxide, carbon monoxide poisoning or manganese intoxication. The product will be manufactured at the group's formulation manufacturing facility in SEZ Ahmedabad, (India).

Carbidopa and Levodopa Tablets USP, 10 mg/100 mg, 25 mg/100 mg, and 25 mg/250 mg had annual sales of USD 75 mn in the United States (IQVIA MAT Dec. 2022).

The group now has 359 approvals and has so far filed over 440* ANDAs since the commencement of the filing process in FY 2003-04.

(*as of 31st December 2022)



For further information please contact: The Corporate Communications Department

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