



Dedicated To Life

January 25, 2024

BSE Limited

P J Towers,
Dalal Street,
Mumbai-400001

Code: 532321

National Stock Exchange of India Limited

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

Code: Zyduslife

Re.: Press Release

Dear Sir / Madam,

Please find enclosed a copy of press release dated January 25, 2024 titled **"Zydus receives Final Approval from the USFDA for Gabapentin Tablets (Once-Daily), 300 mg and 600 mg"**.

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 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 Gabapentin Tablets (Once-Daily), 300 mg and 600 mg

The company is the first to receive the final approval for generic tablets

Ahmedabad, India, January 25, 2024

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 Gabapentin Tablets (Once-Daily), 300 mg and 600 mg (USRLD: Gralise® Tablets). Zydus is the first company to receive final approval for generic Gabapentin Tablets (Once-Daily), 300 mg and 600 mg. Following approval, the product will be launched immediately in the US.

Gabapentin is indicated for the management of Postherpetic Neuralgia (PHN). The product will be manufactured at the group’s formulation manufacturing facility at Moraiya, Ahmedabad. Gabapentin Tablets (Once-Daily), 300 mg and 600 mg had annual sales of USD 85 mn in the United States (IQVIA Nov. 2023).

The group now has 384 approvals and has so far filed over 448* ANDAs since the commencement of the filing process in FY 2003-04.

(*as of 30th September 2023)

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**PRESS
RELEASE**

For further information please contact :
The Corporate Communications Department

Zydus Lifesciences Limited
(formerly known as Cadila Healthcare Limited)

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