



February 18, 2026

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 February 18, 2026, titled **“Zydus receives final approval from USFDA for Bosentan tablets for oral suspension, 32 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.

Yours faithfully,
For, **ZYDUS LIFESCIENCES LIMITED**

DHAVAL N. SONI
COMPANY SECRETARY AND COMPLIANCE OFFICER
MEMBERSHIP NO. FCS7063

Encl.: As above

Zydus Lifesciences 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 USFDA for Bosentan tablets for oral suspension, 32 mg

Ahmedabad, India, 18 February, 2026

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) for Bosentan tablets, oral suspension, 32 mg (USRLD: Tracleer® Tablets for Oral Suspension, 32 mg).

Bosentan 32 mg tablets for oral suspension are indicated for the treatment of Pulmonary Arterial Hypertension (PAH), specifically to improve exercise ability and reduce clinical worsening in children (aged 3 years and older) with idiopathic or congenital PAH. It is a dual endothelin receptor antagonist that lowers high blood pressure in the lungs, typically administered based on body weight.

Bosentan 32 mg tablets will be manufactured at the group’s formulation manufacturing facility at SEZ, Ahmedabad.

Bosentan 32 mg tablets had annual sales of USD 9.3 mn in the United States (IQVIA MAT December 2025).

The group now has 432 approvals and has so far filed 505* ANDAs since the commencement of the filing process in FY 2003-04.

*(*As on 31-Dec-2025)*



**PRESS
RELEASE**

For further information please contact :
The Corporate Communications Department

Zydus Lifesciences Limited

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