



July 4, 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 July 4, 2024 titled **"Zydus receives tentative approval from USFDA for Azilsartan Medoxomil Tablets, 40 mg and 80 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**

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 tentative approval from USFDA for Azilsartan Medoxomil Tablets, 40 mg and 80 mg**

*Ahmedabad, India, 04 July, 2024*

Zydus Lifesciences Limited (including its subsidiaries/ affiliates, hereafter referred to as “Zydus”) has received tentative approval from the United States Food and Drug Administration (USFDA) to market Azilsartan Medoxomil Tablets, 40 mg and 80 mg (USRLD: Edarbi® tablets).

Azilsartan is an angiotensin II receptor blocker (ARB) indicated for the treatment of hypertension to lower blood pressure. Lowering blood pressure reduces the risk of fatal and nonfatal cardiovascular events, primarily strokes and myocardial infarctions. Azilsartan medoxomil tablets may be used either alone or in combination with other antihypertensive agents. The drug will be manufactured at the group’s formulation manufacturing facility in Ahmedabad SEZ - II, India.

Azilsartan Medoxomil Tablets had annual sales of USD 89 mn in the United States (IQVIA MAT March 24).

The group now has 398 approvals and has so far filed over 460\* ANDAs since the commencement of the filing process in FY 2003-04.

*(\*as of 31<sup>st</sup> March 2024)*

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**PRESS  
RELEASE**

For further information please contact :  
**The Corporate Communications Department**

**Zydus Lifesciences Limited**

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