

February 28, 2023

**BSE Limited**

1<sup>st</sup> Floor,  
P J Towers,  
Dalal Street,  
Mumbai-400001

**Code: 532321**

**National Stock Exchange of India Limited**

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

**Code: Zyduslife**

**Re.:** Press Release

Dear Sir / Madam,

Please find enclosed copy of two press releases, both dated February 28, 2023, titled (1) **“Zydus receives final approval from the USFDA for Apixaban Tablets, 2.5 mg and 5 mg”** and (2) **“Zydus receives final approval from the USFDA for Olmesartan Medoxomil and Hydrochlorothiazide Tablets”**.

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



## **Zydus receives final approval from the USFDA for Apixaban Tablets, 2.5 mg and 5 mg**

*Ahmedabad, India, 28 February, 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) for Apixaban Tablets, 2.5 mg and 5 mg (USRLD: Eliquis® Tablets).

Apixaban blocks the activity of certain clotting substances in the blood. It is used to lower the risk of stroke or a blood clot in people with a heart rhythm disorder called atrial fibrillation. It is also used to reduce the risk of forming a blood clot in the legs and lungs of people who have just had hip or knee replacement surgery. It is also used for treatment of blood clots in the veins of legs or lungs, and reduces the risk of them occurring again. The drug will be manufactured at the group’s formulation manufacturing facility at Moraiya, Ahmedabad (India).

Apixaban Tablets, 2.5 mg and 5 mg had annual sales of USD 18,876 mn in the United States (IQVIA MAT Dec. 2022).

The group now has 347 approvals and has so far filed over 440\* ANDAs since the commencement of the filing process in FY 2003-04.

(\*as of 31<sup>st</sup> December 2022)

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**PRESS  
RELEASE**

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](http://www.zyduslife.com)  
CIN : L24230GJ1995PLC025878

## **Zydus receives final approval from the USFDA for Olmesartan Medoxomil and Hydrochlorothiazide Tablets**

*Ahmedabad, India, 28 February, 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) for Olmesartan Medoxomil and Hydrochlorothiazide Tablets, 20 mg/12.5 mg, 40 mg/12.5 mg, and 40 mg/25 mg (USRLD: Benicar HCT<sup>®</sup> Tablets).

Olmesartan Medoxomil and Hydrochlorothiazide, as the name suggests is a combination of two medicines: Olmesartan and Hydrochlorothiazide. Olmesartan is an angiotensin II receptor blocker (sometimes called an ARB blocker) while hydrochlorothiazide is a diuretic (water pill). This combination medicine is used to treat high blood pressure (hypertension). Lowering blood pressure may lower risk of a stroke or heart attack. The drug will be manufactured at the group’s formulation manufacturing facility at Ahmedabad SEZ, India.

Olmesartan Medoxomil and Hydrochlorothiazide Tablets, 20 mg/12.5 mg, 40 mg/12.5 mg, and 40 mg/25 mg had annual sales of USD 41.7 mn in the United States (IQVIA MAT Dec. 2022).

The group now has 346 approvals and has so far filed over 440\* ANDAs since the commencement of the filing process in FY 2003-04.

(\*as of 31<sup>st</sup> December 2022)

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**PRESS  
RELEASE**

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**The Corporate Communications Department**

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