

October 17, 2024

BSE Limited Code: 532321

P J Towers, Dalal Street, <u>Mumbai-400001</u>

National Stock Exchange of India Limited

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

Re.: <u>Press Release</u>

Dear Sir / Madam,

Please find enclosed a copy of press release dated October 17, 2024, titled "Zydus receives final approval from USFDA for Fludrocortisone Acetate Tablets USP, 0.1 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

Code: Zyduslife



## Zydus receives final approval from USFDA for Fludrocortisone Acetate Tablets USP, 0.1 mg

Ahmedabad, India, 17 October, 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 Fludrocortisone Acetate Tablets USP, 0.1 mg (USRLD: Florinef Tablets, 0.1 mg).

Fludrocortisone acetate tablets are indicated as partial replacement therapy for primary and secondary adrenocortical insufficiency in Addison's disease and for the treatment of salt-losing adrenogenital syndrome. Fludrocortisone acetate tablets will be produced at the Group's manufacturing site at Moraiya, Ahmedabad.

Fludrocortisone acetate tablets had annual sales of USD 19.9 mn in the United States (IQVIA MAT July 2024).

The group now has 400 approvals and has so far filed over 465\* ANDAs since the commencement of the filing process in FY 2003-04.

(\*as of 30th June 2024)



For further information please contact: The Corporate Communications Department

## **Zydus Lifesciences Limited**

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