Zydus
dedicated to life

Cadila
Healthcare Limited

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CIN:L24230GJ1995PLC025878

June 28, 2017

BSE Limited
1st Floor, P J Towers,
Dalal Street,
Mumbai – 400 001

National Stock Exchange of India Limited
Exchange Plaza, 5th Floor,
Plot No. C/1, G Block,
Bandra Kurla Complex,
Bandra (East),
Mumbai – 400 051

Re.: **Press Release**

Dear Sir / Madam,

We enclose herewith a copy of press release dated June 28, 2017 titled "Zydus receives final approval from the USFDA for Oxybutynin Chloride Extended-Release Tablets".

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, **CADILA HEALTHCARE LIMITED**

UPEN H. SHAH
COMPANY SECRETARY

Press Release
Press Release

Zydus receives final approval from the USFDA for Oxybutynin Chloride Extended-Release Tablets

Ahmedabad, 28 June 2017

Zydus Cadila has received the final approval from the USFDA to market Oxybutynin Chloride Extended-Release Tablets in the strengths of 5 mg, 10 mg, and 15mg.

The drug is used to treat symptoms of overactive bladder and urinary incontinence (urine leakage) and will be produced at the group's formulation manufacturing facility at Moraiya in Ahmedabad. The estimated sale for Oxybutynin Chloride Extended-Release Tablets is $150.9 million *(Source: IMS Health, IMS National Sales Perspective Audit, MAT April 2017, extracted June 2017)*

The group now has more than 120 approvals and has so far filed over 300 ANDAs since the commencement of the filing process in FY 2003-04.

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For further information please contact :
The Corporate Communications Department

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