Zydus
dedicated to life

Cadila
Healthcare Limited

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CIN:L24230GJ1995PLC025878

September 2, 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 September 2, 2017 titled "Zydus receives final approval from the USFDA for Mycophenolate Mofetil for Injection USP".

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

CADILA HEALTHCARE LIMITED AHMEDABAD

Encl.: As above

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Press Release

Press Release

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Zydus receives final approval from the USFDA for Mycophenolate Mofetil for Injection USP

Ahmedabad, 2 September 2017

Zydus Cadila has received the final approval from the USFDA to market Mycophenolate Mofetil for Injection USP, 500 mg/vial. The drug is indicated for use in combination with other drugs i.e., cyclosporine and corticosteroids for the prophylaxis of organ rejection in patients receiving renal, hepatic or cardiac transplants. The drug will be manufactured at the group's formulations manufacturing facility at Moraiya, Ahmedabad.

The group now has more than 140 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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