Zydus
dedicated to life

Cadila
Healthcare Limited

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July 8, 2017

BSE Limited
1st Floor, P J Towers,
Dalal Street,
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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 July 8, 2017 titled "Zydus receives Final Approval from the USFDA for Doxazosin 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

Zydus receives Final Approval from the USFDA for Doxazosin Tablets

Ahmedabad, 8 July 2017

Zydus Cadila has received the final approval from the USFDA to market Doxazosin Tablets USP, 1 mg, 2 mg, 4 mg, and 8 mg.. The drug is a selective alpha blocker used to treat high blood pressure and urinary retention associated with benign prostatic hyperplasia.

Zydus Cadila has also received the tentative approval from the USFDA to market Lurasidone Hydrochloride Tablets, 20 mg, 40 mg, 60 mg, 80 mg, and 120 mg. The drug is an atypical anti-psychotic used in the treatment of schizophrenia.

Both Doxazosin Tablets and Lurasidone Hydrochloride Tablets will be produced at the group's formulations manufacturing facility at the Pharma SEZ in Ahmedabad.

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 :
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