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CIN:L24230GJ1995PLC025878

November 29, 2017

**BSE Limited**  
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Dalal Street,  
Mumbai – 400 001

**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 – 400 051

Re.: **Press Release**

Dear Sir / Madam,

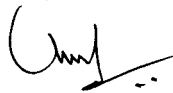
We enclose herewith a copy of press release dated November 29, 2017 titled "Zydus receives final approval from the USFDA for Candesartan Cilexetil and Hydrochlorothiazide Tablets".

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**



Encl.: As above

**Zydus receives final approval from the USFDA for  
Candesartan Cilexetil and Hydrochlorothiazide Tablets**

*Ahmedabad, 29 November, 2017*

Zydus Cadila has received the final approval from the USFDA to market Candesartan Cilexetil and Hydrochlorothiazide Tablets USP in the strengths of 16 mg/12.5 mg, 32 mg/12.5 mg and 32 mg/25 mg. The drug combines an angiotensin II receptor (type AT1) antagonist and a diuretic, hydrochlorothiazide and is used to treat high blood pressure (hypertension). Lowering high blood pressure helps prevent strokes, heart attacks, and kidney problems. It will be manufactured at the group's formulations manufacturing facility at Moraiya, Ahmedabad.

The group now has more than 175 approvals and has so far filed over 310 ANDAs since the commencement of the filing process in FY 2003-04.

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