

24th July 2020

(1) BSE Ltd
Listing Department
Phiroze Jeejeebhoy Towers,
Dalal Street,
Mumbai - 400 001

(2) National Stock Exchange of India Ltd
Listing Department
Exchange Plaza, 5th floor,
Plot no. C/1, G Block,
Bandra Kurla Complex,
Bandra (East), Mumbai - 400 051

Scrip Code: 500087

Scrip Code: CIPLA EQ

(3) SOCIETE DE LA BOURSE DE LUXEMBOURG
Societe Anonyme
35A Boulevard Joseph II,
L-1840 Luxembourg

Sub: Press Release – Cipla receives regulatory approval for launch of Ciplenza (Favipiravir 200 mg) in India to treat mild to moderate COVID-19

Dear Sir / Madam,

Please find enclosed press release dated 24th July 2020 for the captioned subject.

Kindly acknowledge the receipt

This is for your information and records.

Thanking you,

**Yours faithfully,
For Cipla Limited**

**Rajendra Chopra
Company Secretary**

Encl.as above

Prepared by: Juzer Masta



Press Release

Cipla receives regulatory approval for launch of Ciplenza (Favipiravir 200 mg) in India to treat mild to moderate COVID-19

Mumbai, India; July 24, 2020: Cipla Limited (BSE: 500087; NSE: CIPLA EQ, hereinafter referred to as “Cipla”), today announced that it has been granted regulatory approval by the Drug Controller General of India (DCGI) for the launch of Favipiravir in the country under the brand name Ciplenza. The accelerated approval for manufacturing and marketing of the drug is aimed at meeting the urgent and unmet medical need for COVID-19 treatment options in the country through restricted emergency use.

As part of its efforts to enable speedy access to cater to the demand, Cipla will commercially launch Ciplenza in the first week of August priced at Rs 68 per tablet. To ensure fair and equitable distribution of the drug, supplies will be undertaken predominantly through hospital channels and via open channels, prioritised for regions with a high burden of COVID-19 cases.

The drug has been jointly developed by Cipla and CSIR-Indian Institute of Chemical Technology (IICT). As part of this partnership, CSIR-IICT has successfully developed a convenient and cost-effective synthetic process for Favipiravir. The entire process and Active Pharmaceutical Ingredient (API) of the drug has been transferred to Cipla to manufacture and market the drug at scale.

Favipiravir is an off patent, oral anti-viral drug that has been shown to hasten clinical recovery in COVID -19 patients with mild to moderate symptoms.

About Cipla:

Established in 1935, Cipla is a global pharmaceutical company focused on agile and sustainable growth, complex generics, and deepening portfolio in our home markets of India, South Africa, North America, and key regulated and emerging markets. Our strengths in the respiratory, anti-retroviral, urology, cardiology, anti-infective and CNS segments are well-known. Our 46 manufacturing sites around the world produce 50+ dosage forms and 1,500+ products using cutting-edge technology platforms to cater to our 80+ markets. Cipla is ranked 3rd largest in pharma in India (IQVIA MAT June'20), 3rd largest in the pharma private market in South Africa (IQVIA MAT June'20), and is among the most dispensed generic players in the U.S. For over eight decades, making a difference to patients has inspired every aspect of Cipla's work. Our paradigm-changing offer of a triple anti-retroviral therapy in HIV/AIDS at less than a dollar a day in Africa in 2001 is widely acknowledged as having contributed to bringing inclusiveness, accessibility and affordability to the centre of the HIV movement. A responsible corporate citizen, Cipla's humanitarian approach to healthcare in pursuit of its purpose of 'Caring for Life' and deep-rooted community links wherever it is present make it a partner of choice to global health bodies, peers and all stakeholders. For more, please visit www.cipla.com, or click on [Twitter](#), [Facebook](#), [LinkedIn](#).

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