

April 30, 2020

To,  
Dy. General Manager  
Department of Corporate Services,  
BSE Ltd.,  
P. J. Towers, Dalal Street,  
Fort, Mumbai – 400 001.

To,  
The Manager – Listing,  
The National Stock Exchange of India Ltd.,  
Plot No. C/1, G Block,  
Bandra Kurla Complex,  
Bandra (E), Mumbai – 400 051.

Ref: Scrip Code: 532296

Ref: Scrip Name: GLENMARK

Dear Sir,

**Sub: Glenmark receives approval from the regulator (DCGI) to conduct Clinical Trials in India on Favipiravir Antiviral tablets for COVID-19 patients**

With reference to the subject mentioned above, kindly find attached media release which is self-explanatory.

Request you to kindly take the same on record.

Thanking you,

Yours faithfully,  
**For Glenmark Pharmaceuticals Limited**

**Harish Kuber**  
**Company Secretary & Compliance Officer**

**Encl:** as above

Press release

For immediate distribution

## **Glenmark receives approval from the regulator (DCGI) to conduct Clinical Trials in India on Favipiravir Antiviral tablets for COVID-19 patients**

- *Glenmark has successfully developed the API and the formulations for the product through its in-house R&D team*

**Mumbai, India; April 30, 2020:** Glenmark Pharmaceuticals, a research-led, integrated global pharmaceutical company, today announced that it has received approval from the DCGI (Drug Controller General of India), the regulator in India to conduct clinical trials on Favipiravir Antiviral tablets on COVID-19 patients. The product is a generic version of Avigan® of Fujifilm Toyama Chemical Co. Ltd., Japan, a subsidiary of Fujifilm Corporation.

Favipiravir has demonstrated activity against influenza viruses and has been approved in Japan for the treatment of novel influenza virus infections. Recently in the past few months, post the outbreak of COVID-19, multiple clinical trials have been initiated on COVID-19 patients in China, Japan and in the US.

Having internally developed the API and the formulations for the product, Glenmark filed the product for clinical trials with the DCGI and has received approval for conducting the trial on mild to moderate patients. As on date, Glenmark is the first pharmaceutical company in India to be given an approval by the regulator to start the trial on COVID-19 patients in India. As per the clinical trial protocol approved, 150 subjects with mild to moderate COVID-19 will be randomized in the study in a 1:1 ratio to Favipiravir with standard supportive care or standalone standard supportive care. Treatment duration is a maximum of 14 days and the total study duration will be maximum for 28 days from randomization.

***“After having successfully developed the API and the formulations through its in-house R&D team, Glenmark is all geared to immediately begin clinical trials on Favipiravir on COVID-19 patients in India. The clinical trial will let us know the efficacy of this molecule on COVID-19 patients.” said Sushrut Kulkarni, Executive Vice President – Global R&D, Glenmark Pharmaceuticals Limited. He also added, “If the clinical trials are successful, Favipiravir could become a potential treatment for COVID-19 patients.”***

—End—

### **About Glenmark Pharmaceuticals Ltd**

Glenmark Pharmaceuticals Ltd. (GPL) is a global research-led pharmaceutical company with presence across Generics, Specialty and OTC business with operations in over 50 countries. Glenmark’s key therapy focus areas globally are respiratory, dermatology and oncology. It is ranked among the top 80 Pharma & Biotech companies of the world in terms of revenue (SCRIP 100 Rankings published in the year 2019). For more information, visit [www.glenmarkpharma.com](http://www.glenmarkpharma.com)

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**For more information:**

**Glenmark Media Contact**

Madhurima Gupta Jain

Glenmark Pharmaceuticals, Mumbai, India

Tel: +91 22 4018 9606

Email: [corpcomm@glenmarkpharma.com](mailto:corpcomm@glenmarkpharma.com)