

Ref: GLL/SE/2023-24/Dec -

Date: 13/12/2023

To  
The General Manager,  
Corporate Relations Department,  
**BSE Limited**  
Phiroze Jeejeebhoy Towers,  
Dalal Street, **Mumbai - 400001**.  
Maharashtra State, India.  
**Script Code: 531739**

To  
The Listing Manager,  
**The Ahmedabad Stock Exchange Limited**  
A-2, Kamdhenu Complex,  
Opp. Sahajanand College,  
120 Feet Ring Road, Panjara Pol, Ambawadi,  
**Ahmedabad - 380015**.  
Gujarat State, India.  
**Script Code:**

To  
**The Calcutta Stock Exchange Limited,**  
#7, Lyons Range, Murgighata,  
Dalhousie, **Kolkata - 700001**,  
West Bengal State, India.  
**Script Code: 26178**

Dear Sir/Madam,

Sub: Company obtained Good Manufacturing Practices Certificate from  
Drugs Control Administration, Government of Telangana,

Ref: BSE Security ID: GENNEX, Script Code: 531 739

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We are pleased to inform that the Company has received **Good Manufacturing Practices Certificate** from the Drugs Control Administration, Government of Telangana, for a period of ONE year from the date of issue which is valid upto December 10, 2024.

The said GMP Certificate is enclosed with this letter for your information and for taking the same on record.

Thanking you,

Yours faithfully  
**For Gennex Laboratories Limited**

  
**Dinesh Kumar Kejriwal**  
Company Secretary & Compliance Officer  
Membership #A19293



## Gennex Laboratories Limited

**L.Dis.No:125306/TS/2023**

**Dated:12/12/2023**  
**Valid until:10/12/2024**

## **GOOD MANUFACTURING PRACTICES CERTIFICATE**

This is to certify that **M/s GENNEX LABORATORIES LIMITED** situated at address **SY.NO.133,, BOLLARAM VILLAGE, JINNARAM MANDAL, SANGAREDDY DISTRICT,PINCODE 502325,TELANGANA STATE,INDIA** is holding Drug Manufacturing Licence in **Form 25** bearing No. **169/MD/AP/95/B/R** Date.**01/01/2023** Valid upto **31/12/2027** for manufacture for sale or distribution of drugs approved by this Department.The firm is subjected to periodical inspection by this Department.

The firm is following **GOOD MANUFACTURING PRACTICES** as stipulated under the provisions of Schedule "M" of the Drugs and Cosmetics Rules, 1945. The firm should however carry out self inspection from time to time to ensure that the requirements of Good Manufacturing Practices are complied with.

This certificate is valid for one year from the date of issue, unless the firm's manufacturing license is suspended or cancelled by the Licenseing Authority/the firm failed to pay the required the license retention fee.



Digitally Signed By  
**PATLOLLA SARALA**  
Deputy Director and Certifying Authority  
DRUGS CONTROL ADMINISTRATION  
TELANGANA STATE  
Date:12-12-2023 14:02:03 PM

**This Document is Digitally Signed. Signature is not required**