

30-11-2017

Regd.

From

DAO-cum-Licensing Authority
Directorate of AYUSH Haryana,
Sector-3, Panchkula.

To

M/s. La-Medica (India) Pvt. Ltd.,
Plot No. 168, Sector-7,
IMT Manesar, Gurugram.

Memo No. 45/228/Drug-I/AY/HR/2017/ 32763
Dated, Panchkula the 30/11/17

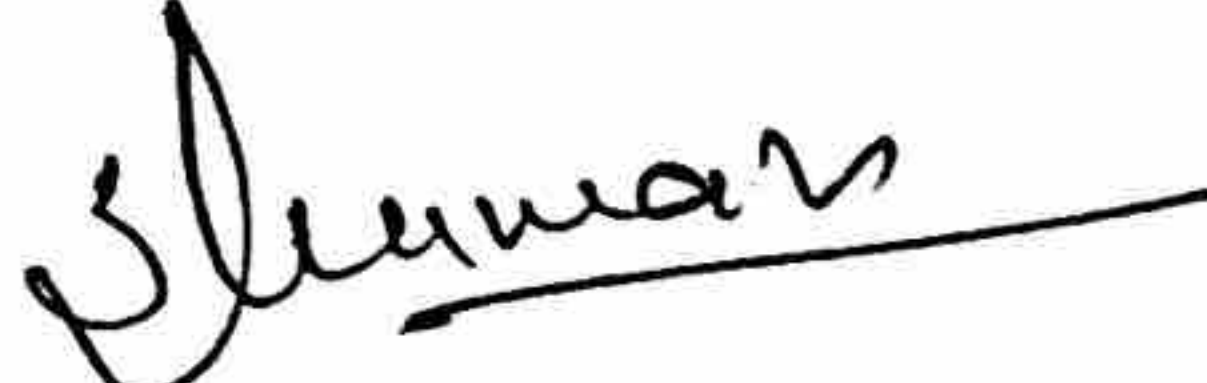
Subject:- Regarding G.M.P. Certificate.

Reference your letter no. LM/609/2017-18/MD/01 dated 05-09-2017 on the subject noted above.

The G.M.P. Certificate is enclosed herewith as desired.

Encl:- One Page.

Endst. No. 45/228/Drug-I/AY/HR/2017/


DAO-cum-Licensing Authority
Directorate of AYUSH Haryana
Dated.

A copy is forwarded to the District Ayurvedic Officer/Drug Inspector, Gurugram with reference to their letter no. 30641 dated 13-11-2017 for information and necessary action.

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DAO-cum-Licensing Authority
Directorate of AYUSH Haryana

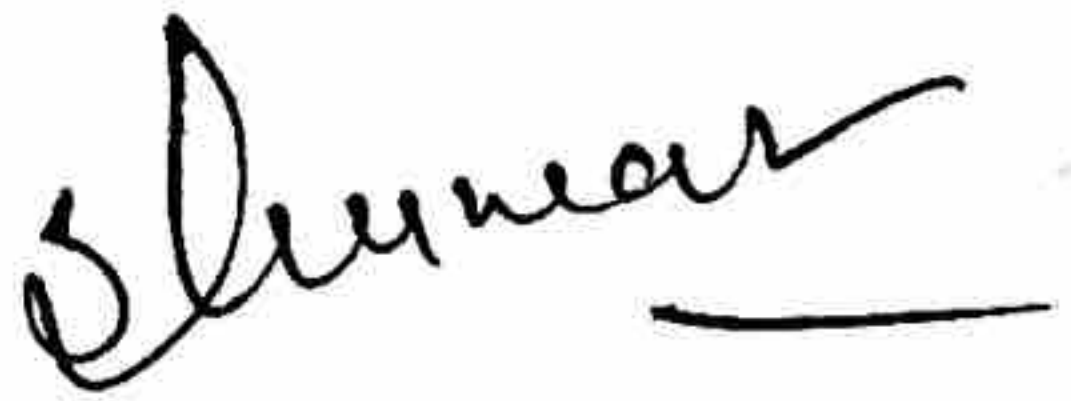
FORM 26-E-1

(See rule 155-B & 157)

CERTIFICATE OF GOOD MANUFACTURING PRACTICES (GMP) TO MANUFACTURER OF AYURVEDA/SIDDHA OR UNANI DRUGS.

Certified that Manufacturing Unit Licensee Namely **M/s. La-Medica (India) Pvt. Ltd., Plot No-168 Sector-7, IMT Manesar, Distt. Gurugram** License No. **13-ISM (HR)** comply with the requirement of Good Manufacturing Practices of Ayurveda/Siddha/Unani Drugs as Laid down in Schedule 'T' of the Drugs and Cosmetic Rules, 1945.

This Certificate is Valid up to **28-10-2020**.


DAO-cum-Licensing Authority,
Directorate of AYUSH Haryana.