DRUGS CONTROL ADMINISTRATION
Government of Telangana

L.Dis.No:80711/TS/2022 **Dated:13/06/2022**
Valid until:11/06/2025

Sub: Drugs and Cosmetics Act, 1940 and Rules made thereunder – Issue of World Health Organisation G.M.P. Certificate– Regarding

Ref: 1. Your letter dated:**28/01/2022.**
2. Joint Inspection report .

-x-x-x-x-

With reference to your application cited, I forward herewith **World Health Organisation GOOD MANUFACTURING PRACTICE** Certificate for the products mentioned in the Joint Inspection Report of the Officers of Drugs Control Administration,Telangana State & Drugs Inspector, CDSCO, Hyderabad vide reference 2nd cited.

Digitally Signed By
PATLOLLA SARALA
Deputy Director and Certifying Authority
DRUGS CONTROL ADMINISTRATION
TELANGANA STATE
Date:13-06-2022 11:23:16 AM

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LIST OF PRODUCTS APPROVED UNDER WHO-GMP CERTIFICATION SCHEME FOR EXPORT PURPOSE

S.No	Generic Name	Brand Name	Composition	PackSize	Market
1	Amisulpride - Ph.Eur				Export
2	Brimonidine Tartrate - Ph.Eur				Export
3	Carvedilol - Ph.Eur / USP				Export
4	Dronedarone Hydrochloride - USP				Export
5	Eszopiclone - USP / IH				Export
6	Itopride Hydrochloride - IH				Export
7	Lanthanum Carbonate - IH				Export

8	Levocetirizine Di hydrochloride - USP / IH				Export
9	Linezolid - USP				Export
10	Racecadotril - Ph.Eur				Export
11	Thalidomide - USP				Export

Manufacturer: M/S **M/s SYMED LABS LIMITED, PLOT NO. 25/B, PHASE-III, IDA, JEEDIMETLA, HYDERABAD, JEEDIMETLA VILLAGE, QUTHBULLAPUR MANDAL, MEDCHAL - MALKAJGIRI DISTRICT,PINCODE 500005,TELANGANA STATE,INDIA**

Drug License No: **26/RR/AP/2003/B/R**,
Dated:28/05/2013 **Under Form 25** ,valid upto **27/05/2023**

When applicable Placing the product on the market as detailed below.

The Unit M/S **M/s SYMED LABS LIMITED, PLOT NO. 25/B, PHASE-III, IDA, JEEDIMETLA, HYDERABAD, JEEDIMETLA VILLAGE, QUTHBULLAPUR MANDAL, MEDCHAL - MALKAJGIRI DISTRICT,PINCODE 500005,TELANGANA STATE,INDIATelangana State,India** was inspected jointly by

It is certified that:

i. The above products had been authorized to be placed on the market for use in the country and exported countries
ii. The manufacturing plant in which the product is produced is subject to inspection at suitable intervals.
iii. The manufacturer conforms to requirements for Good Manufacturing Practices in the manufacture and Quality Control (As recommended by the World Health Organisation) in respect of products to be sold or distributed with in the Country of origin (or to be exported).

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PATLOLLA SARALA
Deputy Director and Certifying Authority
DRUGS CONTROL ADMINISTRATION
TELANGANA STATE
Date:13-06-2022 11:23:16 AM

This Document is Digitally Signed. Signature is not required