DCA

DRUGS CONTROL ADMINISTRATION
Government of Telangana

L.Dis.No.4051/E1/2018 **Dated: 17-12-2018**

To

M/s.Symed Labs Limited,
Plot No.25/B, Phase-III, IDA,
Jeedimetla(V), Quthbullapur(M),
Medchal-Malkajgiri District-500055

Sir,

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: 31.05.2018
2. Joint Inspection report dt: 20.08.2018 & 21.08.2018.

-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 and CDSCO, Zonal Office, Hyderabad vide reference 2nd cited.

This Certificate is valid for Three years from the date of issue.

DRUGS CONTROL ADMINISTRATION HYDERABAD

Yours faithfully,

M.L.V.P.Surendarnath Sai
Joint Director & Licensing Authority(FAC)

INDO LATIN AMERICAN CHAMBER OF COMMERCE
16 APR 2019
NEW-DELHI, INDIA

DCA

DRUGS CONTROL ADMINISTRATION
Government of Telangana

L.Dis.No.4051/E1/2018

Dated: 17-12-2018.

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

1. Amisulpride	IP/Ph.Eur
2. Brimonidine Tartarate	Ph.Eur/IP/IH
3. Carvedilol	Ph.Eur/USP
4. Dronedarone HCL	IH
5. Epalrestat	IH/JP
6. Eszopiclone	IH/USP
7. Itopride HCL	IH
8. Lanthanum Carbonate	IH
9. Levocetirizine Dihydrochloride	IH/IP/USP
10. Linezolid (Form-III)	IP/USP
11. Pregabalin	IH/USP/Ph.Eur
12. Racecadotril	IP/Ph.Eur
13. Thalidomide	USP

Manufacturer : M/s.Symed Labs Limited,
Plot No.25/B, Phase-III, IDA,
Jeedimetla(V), Quthbullapur(M),
Medchal-Malkajgiri District-500055

When applicable : Placing the product on the market as detailed above.

It is certified that the above products had been authorized to be placed on the market for use in the Country.

Drug Licence No. : 26/RR/AP/2003/B/R dated:28.05.2003 under Form – 25 valid upto 27.05.2023

It is also certified that (a) the manufacturing plant in which the product is produced is subject to inspection at suitable intervals.

The Unit M/s.Symed Labs Limited, Plot No.25/B, Phase-III, IDA, Jeedimetla(V), Quthbullapur(M), Medchal-Malkajgiri District-500055 was inspected jointly by Mr.g.Narendra Kumar, DI, CDSCO, Hyderabad and Dr.J.Raju, Drugs Inspector, Jeedimetla(Mfg), Hyderabad on 20.08.2018 & 21.08.2018.

DRUGS CONTROL ADMINISTRATION HYDERABAD GOVERNMENT OF TELANGANA

INDO-AMERICAN CHAMBER OF COMMERCE 16 APR 2019 NEW DELHI, INDIA

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TELANGANA

DRUGS CONTROL ADMINISTRATION
Government of Telangana

L.Dis.No.4051/E1/2018

Dt: 17-12-2018

Issue of WHO GMP CERTIFICATE to M/s.Symed Labs Limited, Plot No.25/B, Phase-III, IDA, Jeedimetla(V), Quthbullapur(M), Medchal-Malkajgiri District-500055

(b) The manufacturer conforms to requirements for Good Manufacturing Practices in the manufacturer and Quality Control (As recommended by the World Health Organisation) in respect of 13 to be sold or distributed with in the Country or origin (or to be exported).

This Certificate is valid for Three years from the date of issue.

HYDERABAD

M.L.V.P.Surendarnath Sai
Joint Director & Licensing Authority(FAC)

LATIN AMERICAN CHAMBER OF COMMERCE INC.
16 APR 2019
NEW-DELHI, INDIA

Attested
SHALINI
Executive Secretary

001449

Coc + Apostille

Attested
SHALINI
Executive Secretary