



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-

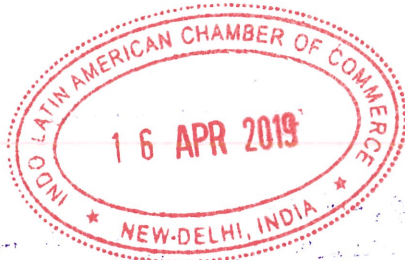
With reference to your application cited, I forward herewith **WORLD HEALTH ORGANISATION GOOD MANUFACTURING PRACTICE** Certificate for the products mentioned in the Joint Inspection Report of the Officers of Drugs Control Administration, Telangana State and CDSCO, Zonal Office, Hyderabad vide reference 2nd cited.

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



Yours faithfully,

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



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.

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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.



[Signature]

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



Attested
[Signature]
SHALINI
Executive Secretary

001449

Coc + Apostille

Attested
SHALINI
Executive Secretary

भारत सरकार GOVERNMENT OF INDIA
अपोस्टिल / APOSTILLE
(Convention de La Haye du 5 octobre 1961)
INDIA

This public document of the type
COMMERCIAL DOCUMENT

is issued to SYMED LABS LTD.

has been signed by SHALINI

with the seal / stamp of EXEC. SECRETARY, INDO LATIN
AMERICAN CHAMBER OF COMMERCE, NEW DELHI

Certified by
Section Officer(OI) MINISTRY OF EXTERNAL AFFAIRS
on 18-Apr-2019 at NEW DELHI, INDIA

with reference no. DLND0009490419

Seal / Stamp

(भवाना शर्मा)
(Bhavana Sharma)
Section Officer (OI)
सी. पी. वी. प्रभाग / C.P.V. Division
विदेश मंत्रालय, नई दिल्ली
Ministry of External Affairs, New Delhi

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MINISTRY OF EXTERNAL AFFAIRS
GOVERNMENT OF INDIA NEW DELHI

acceptance of the Ministry of External Affairs for the documents.