



DRUGS CONTROL ADMINISTRATION
Government of Telangana



L.Dis.No.4052/A3/2018.

Dated 11-10-2018.

To

M/s. Symed Labs Limited, Unit – VI,
Sy.No. 744, 745, 750, 751, 752 & 753,
Mandollagudem Village,
Choutuppal Mandal,
Yadadri District,
Telangana, INDIA.

Sirs,

Sub: Drugs and Cosmetics Act, 1940 and Rules made thereunder – Issue of World Health Organization Good Manufacturing Practice Certificate – Regarding.

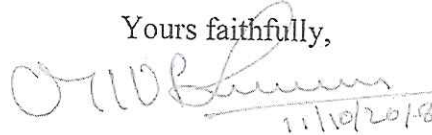
Ref: 1. Your application dated 06.06.2018.
2. Joint Inspection Report dated 09.08.2018 & 10.08.2018.

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I forward herewith **WORLD HEALTH ORGANIZATION GOOD MANUFACTURING PRACTICE CERTIFICATE** for the products recommended by the Joint Inspection Team consisting of officers of Central Drugs Standard Control Organization and officer from Drugs Control Administration, Telangana for **Export** Purpose.

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

Yours faithfully,


11/10/2018

M.L.V.P. SURENDRANATH SAI
JOINT DIRECTOR(FAC)
LICENSING & CONTROLLING AUTHORITY



L.Dis.No.4052/A3/2018.

Dated 11/10/2018.

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

- | | |
|------------------------------|-------------------|
| 1. KETOROLAC TROMETHAMINE | USP/IP/BP/Ph.Eur. |
| 2. LINEZOLID | IP/IH |
| 3. TAPENTADOL HYDROCHLORIDE | IH |
| 4. ONDANSETRON HCL DIHYDRATE | USP |
| 5. PRUCALOPRIDE SUCCINATE | IH |

Manufacturer : M/s. Symed Labs Limited, Unit – VI,
Sy.No. 744, 745, 750, 751, 752 & 753,
Mandollagudem Village, Choutuppal Mandal,
Yadadri District, Telangana, INDIA.

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

It is certified that these products had been authorized to be placed on the market for use in the
country and exporting countries.

Drug Licence No. : 1/YD/TS/2017/B/G, dated 02.01.2017,
in Form – 25

The Unit of M/s. Symed Labs Limited, Unit – VI, situated at Sy.No. 744, 745, 750, 751,
752 & 753, Mandollagudem Village, Choutuppal Mandal, Yadadri District, Telangana, INDIA, was
inspected by Mr. Bikramadithya Chowdhary, Drugs Inspector, O/o the Deputy Drugs Controller(India),
CDSCO, Zonal Office, Hyderabad, Mr. N. Narsiah, Assistant Director, Drugs Control Administration and
Mr. G. Anil Reddy, Drugs Inspector, Drugs Control Administration, Hyderabad on 09.08.2018 &
10.08.2018.

The manufacturer conforms to requirement for Good Manufacturing Practices in the
manufacturing and quality control (As recommended by the World Health Organization) in respect of the
products mentioned above (FIVE in number) for Export in the international market.

This certificate is valid for a period of Three years from the date of issue.




M.L.V.P. SURENDRANATH SAI
JOINT DIRECTOR(FAC)
LICENSING & CONTROLLING AUTHORITY

To
M/s. Symed Labs Limited, Unit – VI,
Sy.No. 744, 745, 750, 751, 752 & 753, Mandollagudem Village,
Choutuppal Mandal, Yadadri District, Telangana, INDIA.




SHALINI
Executive Secretary

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The Ministry of External Affairs
accepts responsibility for the
content of the documents.



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

GOVERNMENT OF 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 02-Nov-2018 at NEW DELHI, INDIA

with reference no. DLDL0033886618

Seal / Stamp



Signature

(DEBABRATA PAUL)
Section Officer (O.I.)
Ministry of External Affairs
New Delhi