

DRUGS CONTROL ADMINISTRATION Government of Telangana



L.Dis.No:90517/TS/2022

Dated:27/08/2022 Valid until:25/08/2025

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: 02/06/2022.

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
RAMDHAN GUGULOTH
Deputy Director and Certifying Authority

DRUGS CONTROL ADMINISTRATION TELANGANA STATE

Date:27-08-2022 12:49:05 PM

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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	DAPSONE - USP/Ph.EUR/BP		######################################		Export
2	FAMOTIDINE - USP/PH.EUR/BP/IP	01		VI	Export
3	FOSPHENYTOIN SODIUM - USP	Car.		10/0	Export
4	RANITIDINE HIDROCHLORIDE - IP/BP/USP/PH.EUR			1	Export
5	ROPIVACAINE HYDROCHLORIDE - USP/Ph.EUR				Export
6	SILDENAFIL CITRATE - IP/USP/PH.EUR/BP				Export



DRUGS CONTROL ADMINISTRATION Government of Telangana



Manufacturer: M/S M/s SMS LIFESCIENCES INDIA LIMITED, UNIT-I,

SY NO. 180/2, KAZIPALLI (V), JINNARAM (M), SANGAREDDY DISTRICT, TELANGANA-502319, INDIA, KHAZIPALLY VILLAGE, JINNARAM MANDAL, SANGAREDDY DISTRICT, PINCODE 502319, TELANGANA STATE, INDIA

Drug License No: 161/MD/AP/96/B/CC,

Dated:01/04/2016 Under Form 25, valid upto 30/03/2026

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

The Unit M/S M/s SMS LIFESCIENCES INDIA LIMITED, UNIT-I, SY NO. 180/2, KAZIPALLI (V), JINNARAM (M), SANGAREDDY DISTRICT, TELANGANA-502319, INDIA, KHAZIPALLY VILLAGE, JINNARAM MANDAL, SANGAREDDY DISTRICT, PINCODE 502319, 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).

Digitally Signed By

RAMDHAN GUGULOTH

Deputy Director and Certifying Authority
DRUGS CONTROL ADMINISTRATION
TELANGANA STATE

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This Document is Digitally Signed. Signature is not required