

L.Dis. No: 164780/TS/2025

Dated :15/02/2025

Valid until:14/02/2028

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: **07/11/2024**.  
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.



*Patolla Sarala*  
15/2/2025

**PATLOLLA SARALA**

Deputy Director and Certifying Authority  
DRUGS CONTROL ADMINISTRATION  
TELANGANA STATE

Date:15-02-2025 13:16:33 PM

L. Dis. No: 164780/TS/2025

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

S. No	Generic Name	Brand Name	Composition	Pack Size	Market
1	CINNARIZINE - BP/EP	NA	NA	NA	Export
2	DIACEREIN - EP	NA	NA	NA	Export
3	SILDENAFIL CITRATE - BP/EP/USP	NA	NA	NA	Export

Manufacturer: M/s RAKSHIT DRUGS PVT LTD,  
SY.NO.10/B, IDA, GADDAPOTHARAM VILLAGE, JINNARAM MANDAL,  
SANGAREDDY DISTRICT, PIN CODE-502 319, TELANGANA STATE, INDIA

Drug License No: 67/MD/AP/2000/B/R,

Dated:01/01/2022 Under Form 25, valid up to 31/12/2026

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







**DRUGS CONTROL ADMINISTRATION**  
**Government of Telangana**



The Unit M/S. RAKSHIT DRUGS PVT LTD, SY.NO.10/B, IDA, GADDAPOTHARAM VILLAGE, JINNARAM MANDAL, SANGAREDDY DISTRICT, PIN CODE – 502 319, TELANGANA STATE, INDIA

Telangana State, India was inspected jointly by

It is certified that:

- The above products had been authorized to be placed on the market for use in the country and exported countries
- The manufacturing plant in which the product is produced is subject to inspection at suitable intervals.
- 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 3 products to be sold or distributed with in the Country of origin (or to be exported).



*15/2/2025*

**PATLOLLA SARALA**

**Deputy Director and Certifying  
Authority**

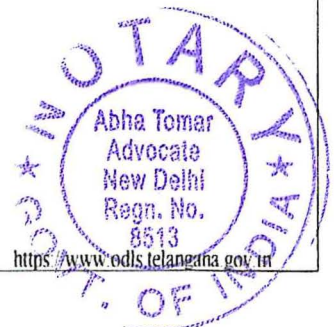
**DRUGS CONTROL ADMINISTRATION  
TELANGANA STATE**

Date:15-02-2025 13:16:33 PM

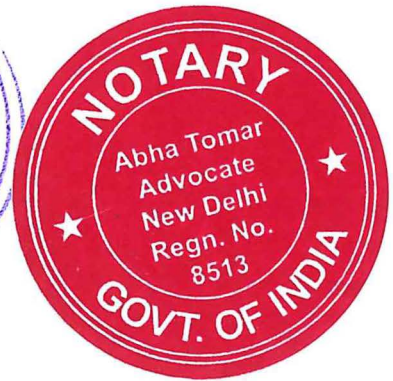
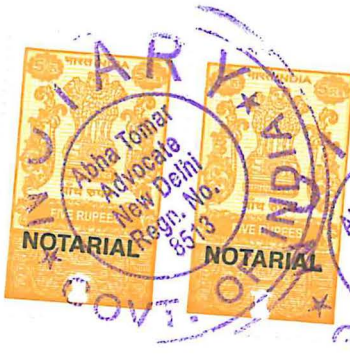
Attested

*Govind Ram*  
**Govind Ram  
Deputy Secretary**

**0 2 8 6 8 9**







ATTESTED

Notary Public, Delhi

5 MAY 2025

The Ministry of External Affairs  
accepts no responsibility for the  
contents of these documents

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

Country **REPUBLIC OF INDIA**

**This public document**  
COMMERCIAL DOCUMENT  
has been signed by **GOVIND RAM**  
acting in the capacity of **DEPUTY SECRETARY**  
bears the seal/stamp of **INDO LATIN AMERICAN CHAMBER OF  
COMMERCE, NEW DELHI**

**Certified**  
at **NEW DELHI, INDIA** the **16-May-2025**  
by **SO (OI/Attestation) MINISTRY OF EXTERNAL AFFAIRS**  
No. **HPHA0000618725**

Seal / Stamp is issued to **RAKSHIT DRUGS PVT LTD**

01 3436988



(मुद्रिका प्रसाद चमार)  
(Mudrika Prasad Chamar)  
अनुभाग अधिकारी (सत्यापन / ओ.आई.)  
Section Officer (Attestation/O.I.)  
सी.पी.वी. प्रभाग / C.P.V. Division  
विदेश मंत्रालय, नई दिल्ली  
Ministry of External Affairs, New Delhi