

DRUGS CONTROL ADMINISTRATION Government of Telangana



L.Dis.No:88919/TS/2022

Dated:05/08/2022 Valid until:03/08/2025

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: **11/05/2022.**

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
B SOWBHAGYA LAXMI
Deputy Director and Certifying Authority

DRUGS CONTROL ADMINISTRATION TELANGANA STATE

Date:05-08-2022 17:07:36 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	ACET <mark>AMIN</mark> OPHEN USP	2	HI O COMPANY	1	Domestic & Export
2	DOMPERIDONE BP/Ph.Eur.	01		1/1	Domestic & Export
3	DOMPERIDONE MALEATE BP/Ph.Eur.	Cast,		فعند ا	Domestic & Export
4	PARACETAMOL BP/Ph.Eur.				Domestic & Export

Manufacturer: M/S M/s SRI KRISHNA PHARMACEUTICALS LIMITED - UNIT - I,

C - 4, INDUSTRIAL AREA, UPPAL, HYDERABAD., UPPAL KHALSA VILLAGE,

UPPAL MANDAL, MEDCHAL - MALKAJGIRI DISTRICT, PINCODE

500039, TELANGANA STATE, INDIA



DRUGS CONTROL ADMINISTRATION Government of Telangana



Drug License No: 30/RR/AP/95/B/R,

Dated:01/01/2013 Under Form 25 ,valid upto 31/12/2022

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

The Unit M/S M/s SRI KRISHNA PHARMACEUTICALS LIMITED - UNIT - I, C - 4, INDUSTRIAL AREA, UPPAL, HYDERABAD., UPPAL KHALSA VILLAGE, UPPAL MANDAL, MEDCHAL - MALKAJGIRI DISTRICT, PINCODE 500039, TELANGANA STATE, INDIA Telangana 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 #Error products to be sold or distributed with in the Country of origin (or to be exported).

Digitally Signed By

B SOWBHAGYA LAXMI

Deputy Director and Certifying Authority

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DRUGS CONTROL ADMINISTRATION
TELANGANA STATE

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This Document is Digitally Signed. Signature is not required