Health & Family Welfare Department
Himachal Pradesh

Certificate of Good Manufacturing Practices

This one page certificate conforms to the format recommended by the **World Health Organization** [General Instructions and Explanatory Notes attached].

Certificate No. HFW-N (ADC) DML/ 2015-16

On the basis of the inspection carried out on 28th & 29th July 2025, we certify that the site indicated on this certificate complies with Good Manufacturing Practices for the dosage forms, categories and activities listed in Table I:

1. Names and Address of Site: **M/s Indo Rama Engineers,**
Plot No. 9, Village Rampur Jattan, Kala Amb,
Distt. Sirmour (H.P.) India.

2. Manufacturer's License No: **N-MNB/16/164** **Form 25**
N-MB/16/165 **Form 28**

3. Table-I:

Dosage Form[s]	Category[ies]	Activity[ies]
Soft Gelatin Capsules	General	Production, Packing & Quality Control

The responsibility for the quality of the individual batches of the pharmaceutical products manufactured through this process lies with the manufacturer.

This certificate now remains valid until **17.09.2028.** It becomes invalid if the activities and/or categories certified herewith are changed or if the site is no longer considered to be in compliance with GMP.

Address of Certifying Authority: **Assistant Drugs Controller**,
O/o State Drugs Controller,
2nd floor, HIMUDA Commercial Complex, Phase-I,
Housing Board, Baddi, Distt. Solan [H.P.] 173205,
INDIA.

Name & Function of Responsible person: **Dr. Kamlesh Naik**
Assistant Drugs Controller
Controlling –cum- Licensing Authority

Telephone/Fax No: 01795-244288, adcbaddi@gmail.com

Date: **18.09.2025**

18 SEP 2025

Signature:
Stamp:

(Dr. Kamlesh Naik)
Assistant Drugs Controller
Cum Licensing Authority
O/o State Drugs Controller
Baddi, Distt. Solan, H.P.173205
adcbaddi@gmail.com, 01795 244288

Assistant Drugs Controller-cum licensing Authority ☆ H.P ☆