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. DCA/SLN/DML/86/10

On the basis of the inspection carried out on 06th & 07th July 2022, 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 Zee Laboratories Limited, Behind 47, Industrial Area, Paonta Sahib, Distt. Sirmour (H.P.) INDIA**

2. Manufacturer's License No: **Form -25: S-MNB/10/67 & Form 28: S-MB/10/68 Form 28D:- N-MB/15/157**

3. Table-I:

Dosage Form[s]	Category[ies]	Activity[ies]
Tablets, Capsules, Oral Liquids, Dry Syrups, Soft Gelatin Capsules, Oral Powder, External Preparations(Creams, Ointments, Gels, Lotions, Shampoo etc.), Small Volume Parenterals Liquid (Vials & Ampoules), Ophthalmic Preparations (Liquid & Eye Ointment), Small Volume Parenterals (Dry Powder Inj)	General	Production, Packing & Quality Control
Tablets, Hard Gelatin Capsules, Oral Powder and Small Volume Parenterals (Dry Only)	Beta Lactum	Production, Packing & Quality Control
Tablets, Capsules, and Small Volume Parenterals (Liquid)	Hormonal	Production, Packing & Quality Control
Tablets, Capsules, and Small Volume Parenterals (Vials & Ampoules) and Small Volume Parenterals (Lyophilized and Dry)	Cytotoxic (Anti Cancer)	Production, Packing & Quality Control
Large Volume Parenterals	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 **10.02.2026.** 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

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

Date: 13.08.2025

Signature:

Stamp: Assistant Drugs Controller-cum licensing Authority H.P.

(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

13 AUG 2025

Explanatory Notes:

1 This certificate, which is in the format recommended by WHO certifies the status of the site, listed in point I of the certificate.

2 The certificate number should be traceable within the regulatory authority issuing the certificate.

3 Where the Regulatory Authority issues a license for the Site, this number should be specified. Record 'Not Applicable" in cases where there is no legal framework for the issuing of a license.

4 Table I

List the Dosage Forms, starting materials, categories and activities. Examples are given below:

Example 1

Pharmaceutical Product[s]1	Category [ies]	Activity [ies]
Dosage Form [s]:		
Tablets	Cytotoxic	Packaging
	Hormone	Production, Packing, Quality Control
	Penicillin	Repackaging and Labeling
Injectables	Cephalosporin	Aseptic preparation, Packaging, Labeling

Example 2

Pharmaceutical Product[s]1	Category [ies]	Activity [ies]
Starting Material [s]		
Paracetamol	Analgesic	Synthesis, Purification, packing, Labeling

Use, whenever available, International Non proprietary Names [Inns] or otherwise national Non proprietary Names

5 The certificate remains valid until the specified date. The certificate becomes invalid if the activities and/or categories certified are changed or if the site is no longer considered to be in compliance with GMP.

6 The requirements for good practices, the manufacture and quality control of drugs referred to in the certificate are those included in Quality Assurance of Pharmaceuticals: a compendium of guidelines and related materials. Good Manufacturing Practices and Inspection. Volume 2, 1999 World Health Organization. Geneva and subsequent updates.