Health & Family Welfare Department
Himachal Pradesh
Baddi, Distt. Solan

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-H [Drugs] 20/18

On the basis of the inspection carried out on **21st & 22nd May 2024**, 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 DSR Life-Care**
At: Village Bersan, PO Lodhimajra
Baddi, Distt. Solan, H.P.-173205, India

2. Manufacturer's License No: L/21/2537/MNB valid upto 19.04.2026
L/21/2538/MB valid upto 19.04.2026

3. Table-I:

Dosage Form[s]	Category[ies]	Activity[ies]
Tablets, Oral Liquid & External Preparations	General	Production, Packing & Quality Control
Tablets & Dry Syrup	Betalactum	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 remains valid until **19.04.2026**. 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**
-cum- Licensing Authority
Himuda Complex, Phase-1, Sai Road,
Baddi, Distt. Solan [H.P.] -173 205
01795-244288,sdc4hp@gmail.com

Name & Function of Responsible person: **Dr. Kamlesh Naik**
Assistant Drugs Controller
-cum- Licensing Authority
H.P., Baddi, Distt. Solan-173205

Date: 19-6-24

Signature: 19/6/24
Stamp: **(Dr. Kamlesh Naik)**
Assistant Drugs Controller
Cum Licensing Authority
O/o State Drugs Controller
Baddi, Distt. Solan, H.P.173205
sdc4hp@gmail.com, 01795-244288

Explanatory Notes:

This certificate, which is in the format recommended by WHO certifies the status of the site, listed in point I of the certificate.
The certificate number should be traceable within the regulatory authority issuing the certificate.

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.

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	Analgesic6	Synthesis, Purification, packing, Labeling

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

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.

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.