

**Health & Family Welfare Department
Himachal Pradesh
Baddi, Dist. 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] 63/08

On the basis of the inspection carried out on 6th & 7th May 2021, 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. Name and address of Site: **M/s Maya Biotech Pvt. Ltd.
Vill -Kondi, PO-Thana,
Baddi, Dist.- Solan (H.P.) India**
2. Manufacturer's License No: **MB/08/725 on Form 28
Valid up to 11.11.2023**
3. Table-I:

Dosage Form[s]	Category [ies]	Activity [ies]
Liquid Injections (SVP)	General	Production, Packing & Quality control
Dry Powder Injections	General	Production, Packing & Quality control
Eye/Ear and Nasal Preparations	General	Production, Packing & Quality control
Dry Powder Injections	Cephalosporin	Production, Packing & Quality control
Metered Dose Inhaler	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 remains valid until **11.11.2023**. 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:

State Drugs Controller
Controlling-cum-Licensing Authority
2nd Floor, Himuda Commercial Complex,
Phase-I Housing Board, Baddi, Distt. Solan,
[H.P]-173205, INDIA

Name & Function of
Responsible person: *

Navneet Marwaha
State Drugs Controller
Controlling-cum-Licensing Authority
01795-244288, sdc4hp@gmail.com

Telephone/Fax No:

Date:

31.5.21



Signature:

Stamp:

(NAVNEET MARWAHA)
State Drugs Controller
Controlling cum Licensing Authority
Baddi Distt. Solan (H. P.)-173205
01795-244288,sdc4hp@gmail.com

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.