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-H [DCA] 98/09

On the basis of the inspection carried out on 30th Nov. & 01st Dec., 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 Beta Drugs Ltd.,**
Kharuni-Lodhimajra Road, Vill. Nandpur
Baddi, Distt. Solan (H.P.) INDIA

2. Manufacturer's License No: **MNB/09/748** **on Form 25**
MB/09/749 **on Form 28**

3. Table-I:

Dosage Form[s]	Category[ies]	Activity[ies]
Tablets	**Oncology**	**Production, Packing & Quality Control**
Capsules	**Oncology**	**Production, Packing & Quality Control**
Small Volume Parenterals	**Oncology**	**Production, Packing & Quality Control**
Lyophilized	**Oncology**	**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 is valid until **23.12.2025.** 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

Telephone/Fax No: 01795-244288, sdc4hp@gmail.com
Date: 24.12.2022

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

24-12-2

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.