Health & Family Welfare Department
Baddi, Distt. Solan, 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[Drugs]231/05(Vol-XII)

On the basis of the inspection carried out on 7th & 8th April - 2021, I 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 Venus Remedies Limited, Hill Top Industrial Estate, Jharmajri, EPIP Phase-I (Extn.), Bhatoli Kalan, Baddi, Distt. Solan, Himachal Pradesh, 173205, India.**

2. Manufacturer's License No: **MB/05/204, & MNB/16/919**

3. Table I. **(a) MB/05/204.**

Dosage Form[s]	Category [ies]	Activity[ies]
Dry Powder Injections	**Cephalosporins**	**Production, Packing & Quality Control**
Dry Powder Injections	**Carbapenems**	**Production, Packing & Quality Control**
Liquid & Lyophilized Injections	**Oncology**	**Production, Packing & Quality Control**
Liquid (PFS, Ampules, & Vial) and Lyophilized Injections	**General**	**Production, Packing & Quality Control**
Oral Sachet	**Cephalosporins**	**Production, Packing & Quality Control**

Table I. (b) MNB/16/919.

Disinfectants	**External Preparation**	**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 from **14.04.2021 to 13.04.2024.** 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 :

Manish Kapoor
Deputy Drugs Controller,
Cum -Licensing Authority,
Baddi, Distt. Solan (H.P.) -173205
01795-244288, ddc4hp@gmail.com

Name & Function of Responsible Person : **Dr. Manish Kapoor**, Deputy Drugs Controller, Cum- Licensing Authority,

Telephone / Fax No: 01795-244288

Dated: 23/4/21

Signature:

Stamp:

DRUGS CONTROLLER-cum-LICENSING AUTHORITY

(Dr. Manish Kapoor) 23/4/2021
DEPUTY DRUGS CONTROLLER
-cum-LICENSING AUTHORITY
O/o STATE DRUGS CONTROLLER
BADDI DISTRICT SOLAN, H.P-173
E mail ddc4hp@gmail.com
Phone 01795-244288

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 specifies. Record 'Not Applicable' in cases where there is no legal framework for 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
	Pencillin	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.