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 [Drugs] 331/05

On the basis of the inspection carried out on 10th & 11th January, 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 Samarth Life Sciences Pvt. Ltd. Vill. Nangal Uperla, Swarghat Road, Nalagarh, Distt. Solan (H.P.) India

2. Manufacturer's License No:

MNB/05/227 & MB/05/228

3. Table-I:

Dosage Form[s]	Category[ies]	Activity[ies]
Tablets, Capsules	Anti Cancer	Production, Packing & Quality Control
Injectables (Dry, Liquid & Lyophilized)	Anti Cancer	Production, Packing & Quality Control
Tablets	Hormone	Production, Packing & Quality Control
Injection (Liquid)	Hormone	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 18/02/2025. 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 Baddi, Distt. Solan, H.P.173205 01795-244288,sdc4hp@gmail.com

Name & Function of Responsible person:

Navneet Marwaha

State Drugs Controller Controlling cum Licensing Authority

01795-244288

Telephone/Fax No: Date: 19/02/2022

Signature:

Controlling cum Lice

sing Author Baddi Distt. Solan (H.

19.2.72

11795-244288 sdc

Stamp:



Explanatory Notes:

- This certificate, which is in the format recommended by WHO certifies the status of the 1.
- The certificate number should be traceable within the regulatory authority issuing the 2. certificate.
- Where the Regulatory Authority issues a license for the Site, this number should be 3. specified. Record 'Not A
- Applicable" in cases where there is no legal framework for the issuing of a license. 4.
- 5. Table I

List the Dosage Forms, starting materials, categories and activities. Examples are given below: Example 1

Example 1		
Pharmaceutical Product[s]1	Category [ies]	Activity [ies]
Dosage Form [s]:		
Tablets	Cytotoxic	Packaging
	Hormone	
Injectables	Penicillin	Production, Packing, Quality Control Repackaging and Labeling Aseptic preparation, Packaging, Labeling
	Cephalosporin	
Example 2		r proparation, rackaging, Labeling

Example 2

Pharmaceutical Product[s]1	Category [ies]	Activity [ies]
Starting Material [s] Paracetamol		
r aracetanioi	Analgesic	Synthesis, Purification, packing,
Use, whenever available,	International Non-r	Labeling

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

- The certificate remains valid until the specified date. The certificate becomes invalid if 6. the activities and/or categories certified are changed or if the site is no longer considered to be in compliance with GMP.
- 7. 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