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] 427/05

On the basis of the inspection carried out on 26th & 27th Sep. 2019, 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. United Biotech (P) Ltd.**
Village Bagbania,
Baddi,-Nalagarh Road
District- Solan (H.P.) 174 101 INDIA

2. Manufacturer's License No: **MNB/05/254 & MB/05/255**
Valid up to 21.02.2021

. Table-I:

Dosage Form[s]	Category[ies]	Activity[ies]
Tablets	General, Betalactum & Oncology	Production, Packing & Quality Control
Capsules	General, Betalactum & Oncology	Production, Packing & Quality Control
Injectable (Liquid & Dry)	General & Oncology	Production, Packing & Quality Control
Dry Syrups	Betalactum	Production, Packing & Quality Control
Liquid Orals	General	Production, Packing & Quality Control
Ointments	General	Production, Packing & Quality Control
Eye/Ear/Nasal Preparations	General	Production, Packing & Quality Control
Injectables (Lyophilsed)	General & Oncology	Production, Packing & Quality Control
Dry Powder Injections	Betalactum	Production, Packing & Quality Control
Dry Powder Injections with Diluents	Cephalosporin	Production, Packing & Quality Control
Soft Gelatin Capsules	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 now remains valid up to **21.02.2021.** 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: **Deputy Drugs Controller,**
Cum Licensing Authority
O/o State Drugs Controller
Baddi Distt. Solan [H.P.] 173 205, INDIA.

Name & Function of Responsible person: **Manish Kapoor**
Deputy Drugs Controller
Cum Licensing Authority

Telephone/Fax No: 01795-244288
Date: 16/10/19

Signature

Stamp: (MANISH KAPOOR) DEPUTY DRUGS CONTROLLER -cum-LICENSING AUTHORITY O/o STATE DRUGS CONTROLLER BADDI, DISTRICT SOLAN, H.P.-173201
E mail ddc4hp@gmail.com
Phone:01795-244288

DEPUTY DRUGS CONTROLLER-cum-LICENSING AUTHORITY ★ H.P. ★ सत्यमेव जयते

UNITED BIOTECH (P) LTD. Certified NEW DELHI INDIA

NOTARY ★ GOVT. OF GUJARAT (INDIA) ★ ARPI K. SHAH AHMEDABAD DIST. GUJARAT STATE REGD. NO 1282/2010 EXP. DT. 9/2/2020

TRUE COPY

ARPI K. SHAH
NOTARY
GOVT. OF GUJARAT

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.

TRUE COPY

ARPI K. SHAH
NOTARY
GOVT. OF GUJARAT

NOTARY (INDIA) ARPI K. SHAH AHMEDABAD DIST. GUJARAT STATE REGD. NO. 1393/2010 DT. 9/2/2020 GUJARAT (INDIA)