Health & Family Welfare Department
Himachal Pradesh
Baddi, Distt. Solan

WHO 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 2nd & 3rd May 2024, we certify that the site indicated on this certificate complies with Good Manufacturing Practices for the dosage forms, categories and activities listed in Table 1:

1. Names and Address of Site: **M/s United Biotech (P) Ltd.**
Village Bagbania,
Baddi-Nalagarh Road,
Distt. Solan (H.P.) 174 101 INDIA

2. Manufacturer's License Nos: **MNB/05/254 & MB/05/255 valid up to 21.02.2026**

3. Table-1:

Dosage Form[s]	Category[ies]	Activity[ies]
Tablets	General, Betalactum & Oncology	Production, Packing & Quality Control
Capsules (Hard Gelatin)	General, & Oncology	Production, Packing & Quality Control
Oral Sachet	General	Production, Packing & Quality Control
Injectables (Liquid & Dry Lyophilized)	General & Oncology	Production, Packing & Quality Control
Liquid Orals	General	Production, Packing & Quality Control
Nasal Preparations	General	Production, Packing & Quality Control
Dry Powder Injections	Betalactum	Production, Packing & Quality Control
Dry Powder Injections	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 remains valid up to **21-02-2026.** 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
Nagar Panchayat Bhawan, Sai Road
Baddi Distt. Solan [H.P.] 173 205
01795244288,sdc4hp@gmail.com

Name & Function of Responsible person: **Dr. Manish Kapoor**
State Drugs Controller
Licensing cum Controlling Authority

Telephone/Fax No: 01795-244288
Date: 09.05.2024

Signature: Manish 9/5/24
Stamp: State Drugs Controller
Controlling cum Licensing Authority
Baddi,Distt.Solan (H.P.) 173205
01795-244288,sdc4hp@gmail.com

Licensing Authority cum Drugs Controller Himachal Pradesh

000776

ATTESTED
K B Lal Saxena
Executive Assistant
PHD Chamber of Commerce and Industry
New Delhi (INDIA)

PHD Chamber of Commerce and Industry
10 MAY 2024
New Delhi-110016

Explanatory Notes:

This certificate, which is in the format recommended by WHO certifies the status of the site, listed in point I of the certificate.

1. The certificate number should be traceable within the regulatory authority issuing the certificate.

2. 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.

3. Table I

List the Dosage [illegible] categories and activities. Examples are giv

Example 1

Pharmaceutical Prod	
Dosage Form [s]:	
Tablets	
	ality Control
	ng
Injectables	ckaging, Labeling

Example 2

Pharmaceutical Pro	
Starting Material [s	
Paracetamol	packing,

Use, whenever [illegible] mes [Inns] or otherwise natio[illegible]

भारत सरकार GOVERNMENT OF INDIA
अपोस्टिल / APOSTILLE
(Convention de La Haye du 5 octobre 1961)
Country REPUBLIC OF INDIA
This public document
COMMERCIAL DOCUMENT
has been signed by MANISH KAPOOR
acting in the capacity of LICENSING AUTHORITY
bears the seal/stamp of PHD CHAMBER OF COMMERCE AND INDUSTRY, NEW DELHI
Certified
at NEW DELHI, INDIA the 10-May-2024
by SO (OI/Attestation) MINISTRY OF EXTERNAL AFFAIRS
No. DLND0001953324
Seal / Stamp
Signature
Is issued to UNITED BIOTECH (P) LTD
OI 2150523
The Ministry of External Affairs accept no responsibility for the contents of the above documents
C.P.V. Division
Ministry of External Affairs, New Delhi

4. 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.

5. 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.