

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 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 21.02.2031**

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 now remains valid up to **08-05-2027**. 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:

Assistant Drugs Controller,
- cum Licensing Authority
2nd Floor, HIMUDA Complex, Phase-I,
Baddi Distt. Solan [H.P.] 173 205
01795-244288,adcbaddi@gmail.com.

007736

ATTESTED
[Signature]
Kamal Bihari Lal Saxena
Executive Assistant
PHD Chamber of Commerce and Industry
New Delhi (INDIA)

Name & Function of Responsible person:

Dr. Kamlesh Naik
Asstt. Drugs Controller
Controlling cum Licensing Authority

Telephone/Fax No:

01795-244288

Date: 21.02.2026

Signature:

Stamp:

(Dr. Kamlesh Naik) 21 FEB 2026

Assistant Drugs Controller
Cum Licensing Authority
C/o State Drugs Controller
Baddi, Distt. Solan, H.P.173205
adcbaddi@gmail.com, 01795 244288



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 Forms, starting materials, categories and activities. Examples are given below:

Example 1

Pharmaceutical Product[s]		
Dosage Form [s]:	Country	REPUBLIC OF INDIA
Tablets	<i>This public document</i>	
	COMMERCIAL DOCUMENT	
	has been signed by KAMAL BIHARI LAL SAXENA	
	acting in the capacity of EXECUTIVE ASSTT	
Injectables	bears the seal/stamp of PHD CHAMBER OF COMMERCE AND INDUSTRY, NEW DELHI	
		Control
		ng, Labeling

Example 2

Pharmaceutical Product[s]		
Starting Material [s]		
Paracetamol	Certified	
	at NEW DELHI, INDIA the 24-Feb-2026	
	by SO (OI/Attestation) MINISTRY OF EXTERNAL AFFAIRS	ng,
	No. DLND0002893426	

Use, whenever available, otherwise national No.

Seal / Stamp

Signature

is issued to UNITED BIOTECH (P) LTD

01 4537682

4. The certificate r 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.

