**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] 18/06 (Vol-IV)**

**On the basis of the inspection carried out on 6th & 7th April, 2017, 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 Associated Biotech,**
**Village Kishanpura, Guru Majra Road,**
**Nalagarh Road, Baddi, Distt. Solan (H.P.)**

2. Manufacturer's License No: **MNB/06/307 Form 25**
**MB/06/308 Form 28**

3. Table-I:

| Dosage Form[s] | Category[ies] | Activity[ies] |
|---|---|---|
| Tablets | Non Betalactum & Cephalosporins | Production, Packing & Quality Control |
| Dry Syrups | Non Betalactum & Cephalosporins | Production, Packing & Quality Control |
| Capsules | Non Betalactum & Cephalosporins | 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 **07.11.2019.** 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
2nd Floor, HIMUDA Complex, Phase-I
Baddi Distt. Solan [H.P.]173205, INDIA.

Name & Function of Responsible person:

**Navneet Marwaha**
State Drugs Controller
Controlling cum Licensing Authority

Telephone/Fax No:

01795-244288, sdc4@gmail.com

Date: 18.04.2019

Signature:

Stamp: (NAVNEET MARWAHA), 18.4.17
State Drugs Controller
Controlling cum Licensing Authority
Baddi Distt.Solan (H. P.)-173205
01795-244288,sdc4hp@gmail.com

**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.