**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] 253/05**

**On the basis of the inspection carried out on 2nd & 3rd September, 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 Torrent Pharmaceuticals Limited,**
**Village Bhud & Makhnu Majra,**
**Tehsil: Baddi-173205,**
**Distt. Solan (H.P.) India**

8. Manufacturer's License No: **MNB/05/183** **FORM 25**
**MB/05/184** **FORM 28**
**Valid upto 21.09.2025**

9. Table-I:

| Dosage Form[s] | Category[ies] | Activity[ies] |
|---|---|---|
| **Tablets** | **General** | **Production, Packing & Quality Control** |
| **Capsules (Hard Gelatin)** | **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 until **16.10.2022.** 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,
H.P., Baddi, Distt. Solan.

Name & Function of Responsible person: **(Dr. Manish Kapoor)**
**Deputy Drugs Controller**,
-cum-Licensing Authority,
O/o State Drugs Controller,
H.P., Baddi, Distt. Solan.

Telephone/Fax No: 01795-244288
Date: 28.09.2020

Signature:
Stamp:

28/9/2020
(Dr. Manish Kapoor)
DEPUTY DRUGS CONTROLLER
-cum-LICENSING AUTHORITY
O/o STATE DRUGS CONTROLLER
BADDI DISTRICT SOLAN, H.P-173205
E mail ddc4hp@gmail.com
Phone 01795-244288

Assistant Drugs Controller HP Baddi Distt. Solan-173205

***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.***