HEATH & 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)57/2016

On the basis of the inspection carried out on 26th & 27th February 2020, 12th & 13th January 2021 and 11th June 2021, 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 Kwality Pharmaceuticals Ltd., Plot No. 1-A, Industrial Area, Raja Ka Bagh, Distt. Kangra 176201, Himachal Pradesh.**
2. Manufacturer's License No: **NNZ/08/40 & BNZ/08/41 on Form 25 & 28 Valid upto 27.12.2025**
3. Table-I:

Dosage Form[s]	Category[ies]	Activity[ies]
Tablets, Capsule, Liquid Injections and Lyophilized Injections	**Cytotoxic Drugs**	**Production, Packing & Quality Control**
Tablets, Capsules & Oral Dry Syrups & SVP Dry	**Cephalosporin**	**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 **15.02.2025.** 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 Commercial Complex, Phase-1.
Housing Board, Baddi, Distt. Solan [H.P.]
173205, INDIA.

Licensing Authority cum Controller ★ Himachal Pradesh ★

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

Telephone/Fax No: 01795-244288, sdc4hp@gmail.com

Date: 21/02/2024

Signature: 21/2/24
Stamp: 21 FEB 2024

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 [res]	Activity [ies]
Dosage Form [s]:		
Tablets	Cytotoxic	Packaging
	Hormone	Production, Packing, Quality Control
	Penicillin	Repackaging and Labeling
lnjectables	Cephalosporin	Aseptic preparation, Packaging, Labeling

Example 2

Pharmaceutical Product[s] I	Category [ies]	Activity [ies]
Starting Material [s]		
Paracetamol	Analgesic	Synthesis, Purification, packing, Labeling

Use, whenever available, lnternational 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.