

**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)57/2016

On the basis of the inspection carried out on 06th & 7th June 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 I:

1. Names and Address of Site: **M/s Kwaliti Pharmaceuticals Ltd.,
Plot No. 1-A, Industrial Area, Raja Ka Bagh,
Distt. Kangra, Himachal Pradesh (India)
-176201.**
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 remains valid until **27.12.2025**. 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: **Assistant Drugs Controller,
O/o State Drugs Controller,
2nd floor, HIMUDA Commercial Complex, Phase-I,
Housing Board, Baddi, Distt. Solan [H.P.] 173205,
INDIA.**

Name & Function of
Responsible person:

Telephone/Fax No:
Date: 27.06.2024



Signature:
Stamp:

Dr. Kamlesh Naik
Assistant Drugs Controller
01795-244288, sd4hp@gmail.com
(Dr. Kamlesh Naik)
Assistant Drugs Controller
Cum Licensing Authority
O/o State Drugs Controller
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