Health & Family Welfare Department
Himachal Pradesh
Baddi, Distt. Solan

Certificate of Good Manufacturing Practices

This one page certificate confirms to the format recommended by the World Health Organization [General Instructions and Explanatory Notes attached].

Certificate No. HFW-H [DRUGS] 665/12

On the basis of the inspection carried out on 2nd & 3th February, 2022, we certify that the site indicated on this certificate complies with Goods Manufacturing Practices for the dosage forms, categories and activities listed in Table I:

1. Names and Address of Site: **M/s Oxalis Labs**
Village Theda, P.O Lodhimajra
Tehsil Baddi Distt. Solan [HP] INDIA -174101

2. Manufacturers License No.: **MNB/11/847** **Form 25**
MB/11/848 **Form 28**

3. Table-I

Dosage Form [s]	Category [ies]	Activity [ies]
Metered Dose Inhaler	General	Production, Packing & Quality Control
External Preparations	General	Production, Packing & Quality Control
Tablets	General	Production, Packing & Quality Control
Sachet (Pellets & Granules)	General	Production, Packing & Quality Control

The responsibility for the quality of the individual batches of the pharmaceuticals products manufactured through this process lies with the manufacturer.

This certificate now remains valid until **09-02-2025**. It become 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 - 1,
Baddi, Distt. Solan [H.P.] 173205, INDIA.
01795-244288, sdc4hp@gmail.com

Licensing Authority cum Drug Controller * Himachal Pradesh *

Name & Function of Responsible Person:

Navneet Marwaha
State Drugs Controller
Controlling -cum- Licensing Authority
01795-244288, sdc4hp@gmail.com

Telephone /Fax No.:
Date: 10 **/02/2022**

Signature:
Stamp:

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

10.2.22

Digitally signed by Moraru Grigore
Date: 2023.10.17 16:32:36 EEST
Reason: MoldSign Signature
Location: Moldova

Explanatory Note:

1. This Certificate, which is in the format recommended by WHO certificate 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 issue of a license.

4. Table I

List the Dosage Forms, starting material, categories and activities.
Examples are given below:

Example 1

Pharmaceuticals Product [s] 1	Category [ies]	Activity [ies]
Dosage Form [s]:		
Tablets	Cytotoxic	Packaging
	Hormone	Production, Packing, Quality Control
	Penicillin	Repackaging and Labeling
Injectable	Cephalosporin	Aseptic preparation, Packaging, Labeling

Example 1

Pharmaceuticals 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 unit the specified data. The certificate become invalid if the activities and /or categories are changed or if the site is no longer considered to be 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. Goods Manufacturing Practices and Inspection. Volume 2, 1999 World Health Organization. Geneva and subsequent updates.