Health & Family Welfare Department
Himachal Pradesh
Baddi, Distt. Solan (H.P.)

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] 995/14

On the basis of the inspection carried out on 28th & 29th Jan. 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. Name and Address of the Site: **M/s Zuvius Lifesciences Private Limited At Plot No. 107 A & 107B, EPIP, Phase-I, Jharmajri, Baddi, Distt.-Solan (HP) INDIA.**

2. Manufacturer's Licence Nos: **L/13/1340/MB Form 28A**
L/13/1339/MNB Form 25A
Valid up to 11.03.2024

Table-I:

Dosage Form[s]	Category[ies]	Activity[ies]
Tablets	Oncology	Production, Packing & Quality Control
Capsules	Oncology	Production, Packing & Quality Control
Injectables	Liquid & Lyophilized (Oncology)	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 **27.02.2022.** 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,
2nd floor, Himuda Complex, Phase-1,
Baddi, Distt.- Solan [H.P.] 173 205, INDIA.

Name & Function of Responsible person: **Manish Kapoor**
Deputy Drugs Controller
-cum Licensing Authority,
O/o State Drugs Controller,
H.P., Baddi, Distt. Solan-173205
01795-244288, ddc4hp@gmail.com

Date: 20 JUL 2021

Signature:

Digitally signed by Chirtoacă Iurie
Date: 2021.12.15 11:46:14 EET
Reason: MoldSign Signature
Location: Moldova

(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

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 licence for the Site, this number should be specified. Record "Not Applicable" in cases where there is no legal framework for the issuing of a licence.

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]i	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 certified 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.