Office of the Controller Food and Drugs Administration

Madhya Pradesh, Idgah Hills, Bhopal (M.P.)-462001
Tel: 0755-2665385, E-mail: cfdamp@rediffmail.com, fdampbhopal@gmail.com

CERTIFICATE OF GOOD MANUFACTURING PRACTICES

This certificate conforms to the format recommended by the world health organization (general instructions an explanatory notes attached)

Certificate No. : 02/2022 **Valid upto** 13 DEC 2025

On the basis of the inspection carried out on 17.11.2022 & 18.11.2022 we certify that the site indicated on this certificate complies with **Good Manufacturing Practices** for the dosage forms, categories and activities listed in table 1.

1. Name & Address of site: **M/s ZUVIUS LIFESCIENCES PVT LTD**
At: Plot No. 58 to 67, Sector B-1,
Umariya-Dungaria, Tehsil-Shahpura,
District-Jabalpur
(M.P.) India - 482003

2. Manufacturer's license No. **28/8/2016 in Form No. 28A**

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3. Table: 1

Dosage Form(s)	Category (ies)	Activity(ies)
Dry Injection	Cytotoxic	Production , Packing , Labeling, Quality Control
Liquid Injection	Cytotoxic	Production , Packing , Labeling, Quality Control

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

4. This certificate remains valid until 13 DEC 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:

Office of the Controller
Food & Drugs Administration
Idgah Hills Bhopal (Madhya Pradesh)
Telephone No.: 0755-2666058
Fax No. : 0755-2665385

Name of authorized person:

Signature:

Shobhit
Dy. Drugs Controller
& Licensing Authority
Food & Drugs Administration
Madhya Pradesh

Stamp and date: 14 DEC 2022

Licensing Authority,
Food & Drugs Administration
Idgah Hills
Bhopal (Madhya Pradesh)

Digitally signed by Dicusar Vladimir
Date: 2023.07.28 13:00:19 EEST
Reason: MoldSign Signature
Location: Moldova

Explanatory notes:

(1) This certificate which is in the format recommended by WHO, certifies the status of the site listed in point 1 of the certificate.

(2) The certification 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 is not legal framework for the issuing of a licence.

(4) Table 1
List the dosage forms, staring materials, categories and activates, Examples are given below.

Example 1

Pharmaceutical Products(S)[2]	Category (ies)	Activity(ies)
Dosage form (s):		
Tablets	Cytotoxic	Packaging
	Hormone	Production , Packaging , Quality Control
	Penicillin	Repackaging and labeling
	Cefalosporin	Aseptic preparation packaging, labeling

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Example 2

Pharmaceutical Products(S)[2]	Category (ies)	Activity(ies)
Starting material (s)[3]		
Paracetamol	Analgesic	Synthesis, purification packing, labeling

Use, whenever available, International Nonproprietary Name (INNs) or otherwise national non proprietary name.

(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 in the manufacturer and quality control of drugs refereed to in the certificate are those included in Quality Assurance of Pharmaceuticals : a compendium of guidelines and related material. Good Manufacturing Practices and Inspection, Volume 2, 1999 World Health Organization, Geneva and Subsequent updates.