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

No. V/WHO-GMP/I-1/2024/ 3856 **Bhopal, Dated:** 11-07-2025

To,

M/s India Phosphate and Allied Industries Pvt. Ltd.
19C-D-E-F & Part of 20A, Industrial Area,
Maxi Road, Ujjain (M.P.)

Sub: - Issue of Model Certificate of WHO-GMP.

Ref: - Your letter dated

***

Please find enclosed herewith the Model Certificate as desired.

Encl: As above

11/7/2025

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

Deputy Drugs Controller cum
State Licensing Authority
Food & Drugs Administration
Madhya Pradesh

OFFICE OF THE CONTROLLER, FOOD AND DRUGS ADMINISTRATION
MADHYA PRADESH

CERTIFICATE OF GOOD MANUFACTURING PRACTICES

This Certificate confirms to the format recommend by the world health organization
(general instructions and explanatory notes attached)

Certificate No.: 07/2025 **Valid Up to: -** 10 JUL 2028

On the basis of the inspection carried out on 07/05/2025 and 08/05/2025 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. India Phosphate and Allied Industries Pvt. Ltd.**
19C-D-E-F & Part of 20A, Industrial Area,
Maxi Road, Ujjain (M.P.)

2. Manufacturing license No. 25/25/2008 in Form No.25

3. Table: 1

Dosage Form(s)	Category (ies)	Activity(ies)
Bulk Drugs	General	Production, Packing, Labeling, Quality Control

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

5. This certificate remains valid until ..... 10 JUL 2028 ..... The Certificate becomes invalid if the activities and *I* or categories certified herewith are changed or if the site is no longer considered to be in compliance with GMP.

6. This certificate is issued as per **WHO TRS No. 908 of 2003.**

Address of Certifying Authority:
Office of the Controller,
Food & Drugs Administration
Idgah Hills Bhopal (Madhya Pradesh)
Telephone No.0755-2666058
Fax No.0755-26665385
E-mail ID-cfdamp@rediffmail.com

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

Name of authorized person: SHOBHIT
Deputy Drugs Controller cum
Licensing Authority
Food & Drugs Administration
Bhopal (M.P.)

Explanatory notes:

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

(8) The certificate number should be traceable within the regulatory authority issuing the Certificate.

(9) 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.

(10) Table 1
List of the dosage forms, starting materials, categories and activities, Examples are given Below,

Example 1

Dosage Form(s)	**Category (ies)**	**Activity(ies)**
Ointment	**General**	**Production, Packing, Labeling, Quality Control**

Example 2

Pharmaceutical Products(S)[2]	**Category (ies)**	**Activity(ies)**
Starting material(s) [3]		
Paracetamol	**Analgesic**	**Production, Packing, Labeling, Quality, Control**

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

(11) 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.

(12) The required for Good Practices in the manufacturer and quality control of drugs referred 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.