OFFICE OF THE CONTROLLER FOOD AND DRUGS ADMINISTRATION MADHYA PRADESH

No. V/WHO-GMP/14/1-485/2019/3793 Bhopal , dated 24/07/2019

CERTIFICATE OF GOOD MANUFACTURING PRACTICES

This one page certificate conforms to the formal recommended by the World Health Organization (General Instruction and Explanatory notes attached)

Certificate No. 04/2006 **Valid up to** 23 JUL 2022

On the basis of the inspection carried out on dated 02.07.2019 & 03.07.2019 we certify that the site indicated on this certificate complies with Good Manufacturing Practices for the API's, categories and activities listed in Table 1.

1. Name and address of site: M/s Ipca Laboratories Limited, P.O. Sejavta Distt. Ratlam (M.P.) -457002, India.

2. Manufacturer's Licence number: 25/35/83 & 28/20 83

3. Table 1:

Dosage Form	Category (ies)	Activity(ies)
Injectable (SVP)	As Per C.O.P.P. Issued	Production packing & Quality Control
Tablets	As Per C.O.P.P. Issued	Production packing & Quality Control
Liquid Oral	As Per C.O.P.P. Issued	Production packing & Quality Control
API (Active Pharmaceuticals Ingredient)	As Per C.O.P.P. Issued	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 remains valid until dated 23 JUL 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 : **Idgah Hills, Bhopal.**

Name and function of responsible person : **Licencing Authority**
Food & Drugs Administration MP
Idgah Hills, Bhopal (M.P.)

Email fda_hotmail.com, Telephone No. 0755-266058 Fax No. 0755-2665385

Signature

SHOBHIT KOSHTA
Licensing Authority
Food & Drugs Administration
Madhya Pradesh

1. This model certificate for GMP is not part of the WHO Certification Schedule on the quality of Pharmaceuticals products moving in international Commerce.

Explanatory notes:

(1) This certificate, which is in the format recommended by WHO certificate 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 issues a licence for the site this number should be specified record 'not applicable' in case where there is no legal framework for the issuing of a licence.
(4) Table 1

List the dosage forms, starting material, categories and activities. Example give below.

Example 1:

Dosage Form	Category (ies)	Activity(ies)
Injectable (SVP)	As Per C.O.P.P. Issued	Production packing & Quality Control
Tablets	As Per C.O.P.P. Issued	Production packing & Quality Control
Liquid Oral	As Per C.O.P.P. Issued	Production packing & Quality Control
API (Active Pharmaceuticals Ingredient)	As Per C.O.P.P. Issued	Production packing & Quality Control

Example 2:

Pharmaceutical Product(s)'	Category (ies)	Activity(ies)
Starting Material(s)3		
N.A	N.A	N.A

2. Pharmaceuticals Products. Any medicine intended for human use or veterinary product administered to food producing animals, presented in its finished dosage for or as a starting material for use in such a dosage form, that is subject to control by pharmaceuticals legislation in both the exporting state and the importing state.
3. Starting Materials: Any substance of a defined quality used in the production of a pharmaceuticals product, but excluding packaging materials.