

Office of the Controller Food and Drugs Administration  
Madhya Pradesh

No. V/28/GMP/E-21/2019/2723

Bhopal, dated 6-6-19

CORRIGENDUM

M/s Erawat Pharma Limited, Plot No. 512, Sector -3, Pithampur, District - Dhar (M.P.) is holding valid Drugs manufacturing Licence No. 28/13/97 in Form 28 which is valid up to 31-12-2022. GMP certificate has been issued to the firm online by this Administration vide this office letter No. DHRGMP201903377 dated 01-05-2019.

Due to technical error in entering the validity of licence of the firm in the application for obtaining GMP Certificate on M.P. Online portal, the validity of licence No. 28/13/97 in Form 28 was wrongly mentioned as 31-12-2017 which may now be read as 31-12-2022.

Please keep this letter with your original licence for inspection by Authority concerned.

To,  
✓ M/s Erawat Pharma Limited,  
Plot No. 512, Sector -3, Pithampur,  
District - Dhar (M.P.)

  
06/06/2019  
State Licensing Authority

Food and Drugs Administration  
Madhya Pradesh

Encl. No. V/28/GMP/E-21/2019

Bhopal, dated

Copy to:

Drugs Inspector, Food and Drugs Administration, Indore (M.P.) for information.

State Licensing Authority  
Food and Drugs Administration  
Madhya Pradesh

**OFFICE OF THE CONTROLLER FOOD AND DRUGS ADMINISTRATION  
MADHYA PRADESH**



No. : DHRGMP201903377

BHOPAL, Dated:01-May-2019

**G.M.P. CERTIFICATE**

It is certified that **ERAWAT PHARMA LIMITED, Plot No. 512 Sector No. 3 Pithampur, DHAR M.P.** State is holding Drug Manufacturing Licence No. **28/13/97** Dated **01-Jan-2013** in Form **28** Valid up to **31-Dec-2017** for manufacturing for sale or distribution of drugs approved by this Department. The firm is subjected to periodical inspection by this department.

The firm is following GOOD MANUFACTURING PRACTICES for all the items permitted to be manufactured under their Drugs manufacturing Licence as stipulated under the provisions of schedule "M" of Drugs and cosmetics Rules.

The Firm should however carry out self inspection from time to time to ensure that the requirements of Good Manufacturing practices are complied with for manufacture of all the permitted items.

This certificate shall remain valid upto to five years from the date of issue or up to the validity of the licences whichever is earlier subject to the following conditions:

- 1) This GMP Certificate is applicable only for manufacturing of such drugs included in the manufacturing licence at any point of time.
- 2) If any deficiency /non-compliance with regard to provisions of Schedule M to Drugs and Cosmetics Rules, 1945 is observed during the validity of this certificate, the action as taken by State Licensing Authority on the licence shall automatically be applicable to this GMP certificate also.
- 3) In Case any drug manufactured by licensee is declared spurious, the action taken by State Licensing Authority on the licence shall automatically be applicable to this GMP certificate also.
- 4) This GMP Certificate shall also cease to be valid if there is any change in the premises or other facilities related to manufacturing of drugs. In such case fresh GMP Certificate shall be required to be obtained by the manufacturer.

To,

**ERAWAT PHARMA LIMITED**  
Plot No. 512 Sector No. 3 Pithampur,  
(Dist)-DHAR, Madhya Pradesh.

Signature valid

Digitally Signed By **RAJNEESH CHOWDHARY**  
(Food And Drugs Administration)

Date : 14-May-2019 18:59:11 IST

**LICENSING AUTHORITY  
FOOD AND DRUGS ADMINISTRATION  
MADHYA PRADESH**

No. : DHRGMP201903377

BHOPAL, Dated: 01-May-2019

Copy forwarded to :

The Drug Inspector Dharmesh Bigoniya ,C/o Dy. Director, Food & Drugs Administration, INDORE

**LICENSING AUTHORITY  
FOOD AND DRUGS ADMINISTRATION  
MADHYA PRADESH**