

OFFICE OF THE CONTROLLER FOOD & DRUGS ADMINISTRATION
MADHYA PRADESH

No. V/25/E - 307 /2020/ 307

Camp Indore, dated 6/7/2020




TO WHOM SO EVER IT MAY CONCERN

M/s Erawat Pharma Ltd. Plot No. 512, Industrial Area, Sector No. 3, Pithampur, District Dhar (M.P.) is holding Drugs Manufacturing Licence No. 25/02(OL)/2018 in Form 25 Valid up to 15.04.2020. The License has deposited requisite online fee for retention of License as per this office record.

In view of above the above mention License is deemed to be valid for a period of 5 years i.e. up to 15/04/2025.

To,

✓ M/s Erawat Pharma Ltd.
Plot No. 512, Industrial Area,
Sector No. 3, Pithampur,
District Dhar (M.P.)


STATE LICENSING AUTHORITY
FOOD & DRUGS ADMINISTRATION
MADHYAPRADESH

**Office of the Controller Food and Drugs Administration
Madhya Pradesh**

No. V/G/25/E-3/2018

5564

Bhopal, dated

2-11-18

To,

M/s Erawat Pharma Limited
Plot No. 512, Industrial Area, Sector No. III,
Pithampur, District - Dhar -454774 (M.P.)

Sub: Formal Grant of licence in Form 25.

On the above subject it is stated that your firm are already having valid Drugs Manufacturing Licence in Form 28 for manufacturing of Hard Gelatin Capsules Shells IP. Previously due to some technical reasons inclusion permission for Empty Hydroxy Propyl Methyl Cellulose Capsule Shells was granted to the you under Drugs Manufacturing Licence No. 28/13/97 as applied by you.

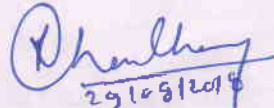
At the time of renewal of Drugs Manufacturing Licence No. 28/13/97, it was noticed that Empty Hydroxy Propyl Methyl Cellulose Capsule Shells comes under the category of non biological products and same was informed to you. You have applied for grant of licence in Form 25 to manufacture Empty Hydroxy Propyl Methyl Cellulose Capsule Shells & submitted the requisite fee.

Therefore, as per your request & fee deposited by you please find enclosed herewith the formal Drugs Manufacturing Licence in Form 25 No. 25/02(OL)/2018 in Form No. 25 for the period of w.e.f. 16-04-2015. The validity of licence shall be 16-04-2015 to 15-04-2020.

Encl: Original licence

Total items: 01 item.

Total pages- 01



29/10/2018
State Licensing Authority
Food and Drugs Administration
Madhya Pradesh

Endt. No. V/G/25/E-3/2018

Bhopal, dated

Copy to:

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


State Licensing Authority
Food and Drugs Administration
Madhya Pradesh

FORM -25

(See Rule 70)

[LICENCE TO MANUFACTURE FOR SALE OR FOR DISTRIBUTION OF] DRUGS OTHER THAN THOSE SPECIFIED IN [SCHEDULE C & C (1) AND X]Number of Licence No. **25/02(OL)/2018**Date of Issue - **29-08-2018**

1. **Shri Abhishek Jain S/o Shri Hemchand Jain, Director of M/s Erawat Pharma Limited** is hereby licensed to manufacture the following categories of drugs being drugs other than those specified in Schedule C & C (1) and X to the Drugs and Cosmetics Rules, 1945 on the premises situated at **Plot No. 512, Industrial Area, Sector No. III, Pithampur, District - Dhar - 454774 (M.P.)** under the direction and supervision of the following competent technical staff.

a) Competent Technical Staff (Names)

1. On Manufacturing Side:

Shri Rakesh Sharma**MC - 1582**

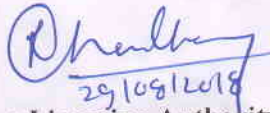
On Testing Side:

Shri R.K. Vyas**ANACHEM- 1205****Shri Shiv Prasad Prajapati****ANACHEM- 23/2016****Ms. Vandana Yadav****ANACHEM- 24/2016**

- b) Names of Drugs (each item to be separately specified) :- **"EMPTY HYDROXY PROPYL METHYL CELLULOSE CAPSULE SHELLS"**

2. The licensee authorises the sale by way of wholesale dealing and storage for sale by the licensee of the drugs manufactured under the licence, subject to the conditions applicable to licence for sale.
3. The licence shall be in force from w.e.f. **16-04-2015** to **15-04-2020**
4. The licence is subject to the conditions stated overleaf and to such other conditions as may be specified in the Rules for the time being in force under the Drugs and Cosmetics Act, 1940.

Date of issue: **29-08-2018**


29/08/2018
State Licensing Authority
Food and Drugs Administration
Madhya Pradesh

CONDITIONS OF LICENCE

1. This licence and any certificate of renewal in force shall be kept on the approved premises and shall be produced at the request of an Inspector appointed under the Drugs and Cosmetics Act, 1940.
2. Any change in the [competent technical staff] named in the licence shall be forthwith reported to the Licensing Authority.
3. If the licensee wants to manufacture for sale additional items of drugs not included above he should apply to the Licensing Authority for the necessary endorsement as provided in Rule 69 (5). This licence will be deemed to extend to the categories so endorsed.
4. The licensee shall inform the Licensing Authority in writing in the event of any change in the constitution of the firm operating under the licence. Where any change in the constitution of the firm takes place, the current licence shall be deemed to be valid for a maximum period of three months from the date on which the change takes place unless, in the meantime, a fresh licence has been taken from the Licensing Authority in the name of the firm with the changed constitution.