FORM 25

(See rule 70)

[LICENCE TO MANUFACTURE FOR SALE OR FOR DISTRIBUTION OF] DRUGS OTHER THAN THOSE SPECIFIED IN [SCHEDULES C, C(1) AND X]

1. Number of License: **HP-MNB-00264** Date of Issue: **27/06/2025** Valid Up To: **26/06/2030** Firm Number: **418**

2. **MK HEALTHCARE** is hereby licensed to manufacture the following categories of drugs being drugs other than those specified in [Schedules C, C(1) and X] to the Drugs and Cosmetics Rules, 1945, on the premises situated at **Khasra No. 1112, Gahlian Khas, Kangra, Himachal Pradesh-176029.**, **Kangra**, **Kangra** under the direction and supervision of the following [competent technical staff]

(a) [Competent technical staff] (Names)

Manufacturing Chemist		
Name	**Qualification**	**Approved in Section(s)**
Anuj Kumar	Bachelor of Science	Tablets, Capsules, Oral Liquid
Analytical Chemist		
Name	**Qualification**	**Approved in Section(s)**
Devendra Kumar Sharma	Bachelor of Science	Chemical Testing, Instrumentation

(b) Names of drugs (each item to be separately specified) : **Oral Solid Dosage Form General Category(Tablets and Capsule)**

3. The licence authorises the sale by way of wholesale dealing and storage for sale by the licensee of the drugs manufactured under the license, subject to the conditions applicable to license for sale.

4. The licence, unless sooner suspended or cancelled, shall remain valid perpetually. However, the compliance with the condition of licence and the provisions of the Drugs & Cosmetics Act 1940 (23 of 1940) and the Drugs Rules 1945 shall be assessed not less than once in three years or as needed per risk-based approach.

5. The licence is subject to the conditions stated below and to such other conditions as may be specified in the Rules for the time being in force under the Drugs & Cosmetics Act 1940.

Date **27/06/2025**

DRUGS LICENSING AUTHORITY (H.P.)

Nishant Sareen

Licensing Authority

H.Q. KANGRA AT DHARAMSHALA HP

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 expert 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.

FORM 28

(See rule 76)

[LICENCE TO MANUFACTURE FOR SALE OR FOR DISTRIBUTION OF DRUGS SPECIFIED IN SCHEDULES C AND C(1) EXCLUDING THOSE SPECIFIED IN SCHEDULES X]

1. Number of License: **HP-MB-00265** Date of Issue: **27/06/2025** Valid Up To: **26/06/2030** Firm Number: **418**

2. **MK HEALTHCARE** is hereby licensed to manufacture at the premises situated at **Khasra No. 1112, Gahlian Khas, Kangra, Himachal Pradesh-176029.**, **Kangra**, **Kangra** the following drugs being drugs specified in Schedules C and C(1) [excluding those specified in Schedules X] to the Drugs and Cosmetics Rules 1945.

(a)Names of Drugs : **Oral Solid Dosage Form General Category(Tablets and Capsule)**

(b) Name of approved [Competent technical staff]. (Names)

Manufacturing Chemist		
Name	**Qualification**	**Approved in Section(s)**
Anuj Kumar	Bachelor of Science	Tablets, Capsules, Oral Liquid
Analytical Chemist		
Name	**Qualification**	**Approved in Section(s)**
Devendra Kumar Sharma	Bachelor of Science	Chemical Testing, Instrumentation

3. The licence 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 licences for sale.

4. The licence, unless sooner suspended or cancelled, shall remain valid perpetually. However, the compliance with the condition of licence and the provisions of the Drugs & Cosmetics Act 1940 (23 of 1940) and the Drugs Rules 1945 shall be assessed not less than once in three years or as needed per risk-based approach.

5. The licence is subject to the conditions stated below and to such other conditions as may be specified in the Rules for the time being in force under the Drugs & Cosmetics Act 1940.

Date **27/06/2025**

DRUGS LICENSING AUTHORITY (H.P.)

Nishant Sareen

Licensing Authority

H.Q. KANGRA AT DHARAMSHALA HP

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. If the licensee wants to undertake during the currency of the licence the manufacture of any drug specified in Schedules C and C (1) [excluding those specified in Schedule X] not included above, he should apply to the Licensing Authority for the necessary endorsement as provided in rule 75(3). This licence will be deemed to extend to the items so endorsed.

3. Any change in the [competent technical staff] shall be forthwith reported to the Licensing Authority.

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.