हिमाचल प्रदेश

DEPARTMENT OF AYURVED
GOVERNMENT OF HIMACHAL PRADESH
Form 26 E-I
Rule 157 (1A)
(CERTIFICATE OF GOOD MANUFACTURING PRACTICES (GMP) TO MANUFACTURER OF AYURVEDA DRUGS)

Certified that manufacturing Unit licensee namely M/S Ayush Herbals & Healthcare Unit-II, Kh.No. 229/103/2, Village Nagal Suketi Road Kala-Amb, Tehsil Nahan, Distt. Sirmour, H.P. 173030 within the state of Himachal Pradesh having Licence No.HP-308-Ay comply with the requirements of Good Manufacturing Practices (G.M.P) of Ayurvedic drugs as laid down in Schedule "T" of the Drugs and Cosmetics Rules, 1945.

This certificate is valid for a period of five years from 01.9.2020 to 31.8.2025 and the Good Manufacturing Practices (GMP) is valid for the various dosage forms or Rasaushadhis, as follows:-
(For Aushadh Ghana (Medicinal Plants Extracts-Dry/Wet) Section Only.

File No. : Ay. H. (A) (3)-308/2020- 16930
Dated: - 03/10/2020
Place: Shimla

Himachal Pradesh Drug Controller Cum Licencing Authority (Ayurveda)

Director Ayurveda-cum
Licensing Authority, Hdqrs.
Licensing Authority (H.P.)

Registered

Corrigendum

DEPARTMENT OF AYURVEDA
HIMACHAL PRADESH
***

No. Ay. H(A)(3)-308/20- 16929 **Dated** 03/10/2020 **Shimla-171009, the**

To

M/S Ayush Herbals & Healthcare Unit-II,
Kh.No. 229/103/2, Village Nagal Suketi Road Kala-Amb
Tehsil Nahan, Distt. Sirmour, H.P. 173030

Subject: - Revised Ayurvedic Drug Manufacturing Licence No. HP-308-Ay and GMP Certificate.

Sir,

With reference to your letter dated 12-09-2020 on the matter cited as subject.

Enclosed please find herewith the revised Ayurvedic Drug Manufacturing Licence No. HP-308-Ay & GMP Certificate.

Encl. As above

Director Ayurveda-Cum-
Licensing Authority, H.P.

DEPARTMENT OF AYURVEDA
HIMACHAL PRADESH, SHIMLA-171009 03/10/2020

No.Ay.H (A)(3)-308 /20 - 16930 **Dated,** Shimla-9, the

To

M/S Ayush Herbals & Healthcare Unit-II,
Kh.No. 229/103/2, Village Nagal Suketi Road Kala-Amb
Tehsil Nahan, Distt. Sirmour, H.P. 173030

Subject: - **Grant of License for the manufacturing of Ayurvedic Medicines.**

Dear Sir,

Reference your application on the above-mentioned subject.

Enclosed herewith please find license No.HP-308-Ay on Form No. 25-D prescribed under the Drugs and Cosmetics Act/Rules, 1945 for the manufacturing of Ayurvedic Medicines.

This license is valid from the date of issue to **31.8.2025.**

The drug manufactured should be labeled as required under Rule 161 of the Drugs and Cosmetics Act/Rules, 1945, provision of Rules 158 A and Schedule T should be strictly adhered to.

If on the inspection it is observed that the conditions of the License and the provisions of the Drugs & Cosmetics Act, 1940 and the Rules made thereunder are not being complied with, the license shall be cancelled and necessary legal action will be taken against you.

Kindly acknowledge receipt of this letter and license.

Yours faithfully,

Director of Ayurveda-cum-
Licensing Authority, H.P

FORM No. 25-D
(See Rule -154)

License to manufacture for sale of Ayurvedic Drugs.

No. of License **HP-308-AY.**

1. **M/S Ayush Herbals & Healthcare Unit-II** is/are hereby licensed to manufacture the following Ayurvedic medicines on the premises situated at **Kh.No. 229/103/2, Village Nagal Suketi Road Kala-Amb, Tehsil Nahan, Distt. Sirmour,** H.P. 173030 under the direction and supervision on the following technical staff:-

Technical Staff (Name)

1. **Dr. Nitika, B.A.M.S (QC)**
2. **Miss, Priya Sharma B.Sc. (P)**
3. **Miss Jyoti Devi, B.Sc. (QC)**
4. **Mr. Gobind Sharma, B.Sc (QC)**

Approval of drugs will be conveyed lateron.

For "Aushadh Ghana" (Medicinal Plants Extracts-Dry/Wet) section only

2. The license shall be in force from **01.9.2020 to 31.8.2025.**
3. The license 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 and Cosmetics Act, 1940.

Date of Issue 03-10-2020

Signature Director Ayurveda -cum-
Designation Licensing Authority (H.P.)

Conditions of License

1. The license 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 changes in the technical staff named in the license shall be reported forthwith to the Licensing Authority.
3. This license shall be deemed to extend to such additional items as the licensee may intimate to the Licensing Authority from time to time, and as may be endorsed by 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 license. Where any change in the constitution of the Firm takes place, the current license 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 license has been taken from the Licensing Authority in the name of the firm with the changed constitution.