Government of Uttar Pradesh
Head Quarter, Food Safety And Drug Administration, Sector C, Aliganj, Lucknow, Lucknow, Lucknow, Uttar Pradesh-226024, India

Licence Retention Letter

To
AOS PRODUCTS PVT LTD
S-33, South Side, G T Road, Industrial Area, Ghaziabad,
Ghaziabad Sadar, District-Ghaziabad, State-Uttar Pradesh,
India -201009

Retention No: RLT2025UP02074
Date: 16-Jun-2025

Subject : - **Drugs & Cosmetic Act- 1940 & rules there under**

Sir,
Please refer to your application number **UP/ML/F24/2025/00023** dated **25-Feb-2025**, we have to inform you that your said application is approved and below mentioned licence is retained, whose validity date is mentioned below :-

New Licence No	Old Licence No	Issue Date	Retention Date	Valid Upto
MLF252025UP000007	01 of 2015	13-Jun-2025	23-Jan-2025	22-Jan-2030

Retained Product list with this Retention application (See **Annexure 'A'**)
Attached Technical staff (See **Annexure 'B')**

Conditions of Licence

1. Subject to no change in previous constituation and already approved premises and validity of Technical manpower.
2. The licensee shall not claim any equities or rights in the property under the refrence on the strength of this Retention Letter.
3. You are required to apply for the Retention of the above licences 3 months before the validity expires.

Date: 16-Jun-2025

Mr. Shashi Mohan Gupta
Licensing Authority

Licence Retention Letter

To
AOS PRODUCTS PVT LTD
S-33, South Side, G T Road, Industrial Area, Ghaziabad,
Ghaziabad Sadar, District-Ghaziabad, State-Uttar Pradesh,
India -201009

Annexure 'B'(List of Technical Staff)

Competent Technical Staff:

Member ID	Member Name	Assign Role / Designation	Qualification
20241859472	Dr. Madhu Gautam	Analytical Chemist	Pg Botany
20245799199	Mr. Inderpreet Singh	Manufacturing Chemist	B.E

प्रेषक,

आयुक्त,
खाद्य सुरक्षा एवं औषधि प्रशासन, उ0प्र0,
9, जगत नारायण रोड, लखनऊ।

सेवा में,

मेसर्स एओएस प्रोडक्ट्स प्राइवेट लिमिटेड,
एस–33, साउथ साइड, जीटी रोड,
इण्डस्ट्रियल एरिया,
गाजियाबाद, उ0प्र0।

सं0: ड्रग/7108/907 लखनऊ दिनांक 30/01/2015

विषय : फर्म को प्रपत्र–25 पर नवीन औषधि निर्माण लाइसेंस निर्गत किये जाने के सम्बन्ध में।

महोदय,

उपर्युक्त विषयक अपने आवेदन पत्र दिनांक 31.01.2014, जिसके द्वारा फार्म–25 पर नवीन औषधि निर्माण लाइसेंस की स्वीकृति का अनुरोध किया गया है, को संदर्भित करें।

आपके आवेदन पत्र पर विचारोपरान्त फार्म–25 पर औषधि निर्माण लाइसेंस संख्या– 01 ऑफ 2015, औषधि एवं प्रसाधन सामग्री अधिनियम 1940 एवं नियमावली 1945 में निहित प्राविधानों के अनुसार स्वीकृत कर संलग्न कर प्रेषित किया जा रहा है। संलग्न औषधि निर्माण लाइसेंस दिनांक 23.01.2015 से 22.01.2020 तक की अवधि हेतु मान्य होगा। औषधि निर्माण लाइसेंस की शर्तों का पालन करना अनिवार्य होगा। उक्त औषधि निर्माण लाइसेंस पर निर्माण किये जाने वाली औषधियो को लाइसेंस पर अंकित कर तथा मानचित्र की अनुमोदित एवं हस्ताक्षरित प्रति भी इस पत्र के साथ संलग्न कर प्रेषित की जा रही है।

संलग्नक–उपरोक्तानुसार।

भवदीय

Malla 23.1.2015

(ए0के0 मलहोत्रा)
औषधि अनुज्ञापन एवं नियंत्रण प्राधिकारी,
उत्तर प्रदेश।

सं0: ड्रग/7108/ तद्दिनांक

प्रतिलिपि–निम्नलिखित को सूचनार्थ एवं आवश्यक कार्यवाही हेतु प्रेषित–

(1) सहायक आयुक्त (औषधि) कार्यालय आयुक्त, मेरठ मण्डल मेरठ।
(2) औषधि निरीक्षक, कार्यालय जिलाधिकारी, गाजियाबाद को लाइसेंस की प्रति एंव निर्माण इकाई के मानचित्र की प्रति सहित।

(ए0के0 मलहोत्रा)
औषधि अनुज्ञापन एवं नियंत्रण प्राधिकारी,
उत्तर प्रदेश।

FORM-25

(See Rules 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 and date of issue: 01 **of 2015;** **Dated:** 23·01·2015

1. **M/s AOS Products Private Limited,** is hereby licenced 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 **S-33, South Side GT Road Industrial Area, Ghaziabad, (U.P.)** under the direction and supervision of the following competent technical staff:

(a) ***Competent Technical Staff (Names) :***

Manufacturing Chemist : Mr. Inderpreet Singh

Analytical Chemist : Mr. Vinod Kumar

(b) **Names of drugs** (each item separately specified): Categories of drugs approved:

Menthol IP/USP, Peppermint Oil IP/USP/BP, Mentha Oil IP, Levo Menthol BP, Dementholised Mint Oil BP

2. 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 licence for sale.
3. The licence shall be in force from 23·01·2015 to 22·01·2020
4. 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 and Cosmetics Act, 1940.

Dated: 23·01·2015

23.01.2015

(A.K. Malhotra)
Drug Licensing Cum Controlling Authority
Uttar Pradesh

(A. K. MALHOTRA)
Drugs Licensing and Controlling Authority
Uttar Pradesh (India)

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. (Omitted vide Notification No. S.O. 289; 20.12.1972).
5. 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.