

FOOD SAFETY & DRUG ADMINISTRATION AUTHORITY
Directorate General of Medical Health & Family Welfare,
Sahastradhara Road, Dehradun, Uttarakhand (INDIA)

File No. : 17P/1/16/2012/ 8603

Dated : 05 / 05 / 2022

LICENSE RETENTION CERTIFICATE

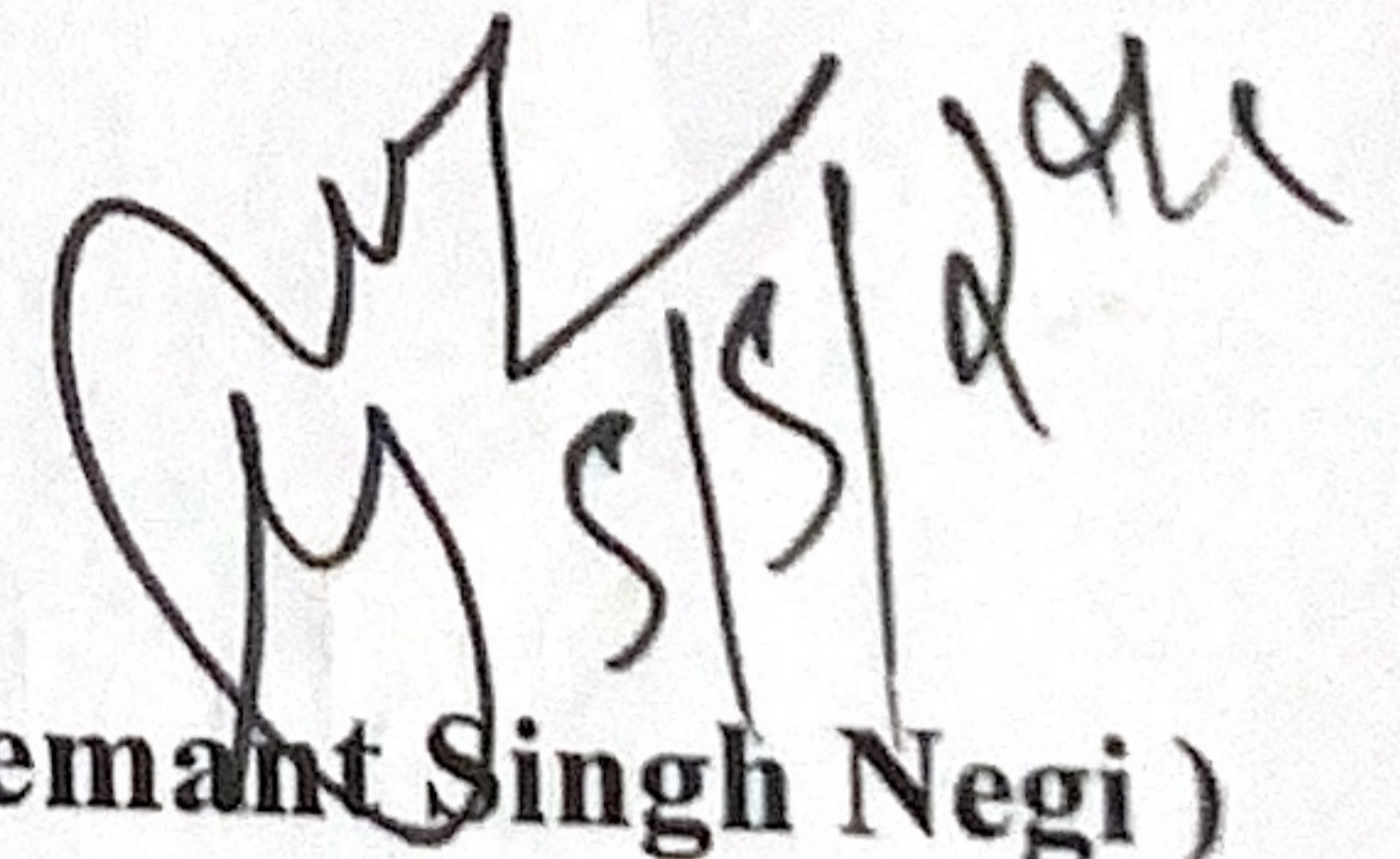
This is to certify that M/s Aryan Biotech Pvt.Ltd., Plot No.16, Sector-6B, IIE, SIDCUL, Ranipur, Distt. Haridwar, Uttarakhand is holding manufacturing license No. 18/UA/2012 on Form-25 & 17/UA/SC/P-2012 on Form-28 valid up to 18-04-2022.

It is further certified that the firm has Submitted necessary license retention fee of Rs 15000/- (Rs fifteen thousand only) vide challan no 883070 Dated 04-04-2022 under the provision of Drug & Cosmetics Rules as amended.

Therefore the above Said license granted to the firm shall deemed to be valid and shall be in force for another 5 years From- 19-04-2022 to 18-04-2027.

Unless it is Suspended or Cancelled During this Period.

This Certificate is issued on the request of the firm.



(Hemant Singh Negi)
Drug Controlling & Licensing
Authority (Mfg.)
Garhwal Mandal.Uttarakhand
(Uttarakhand)

(Hemant Singh Negi)
Drug Controlling & Licensing Authority
(Mfg) Garhwal Mandal
Uttarakhand



FOOD SAFETY & DRUG ADMINISTRATION AUTHORITY
Directorate General of Medical Health & Family Welfare,
Sahastradhara Road, Dehradun, Uttarakhand (INDIA)

File No. : 17P/1/16/2012/ 86-3

Dated : 05 / 05 / 2022

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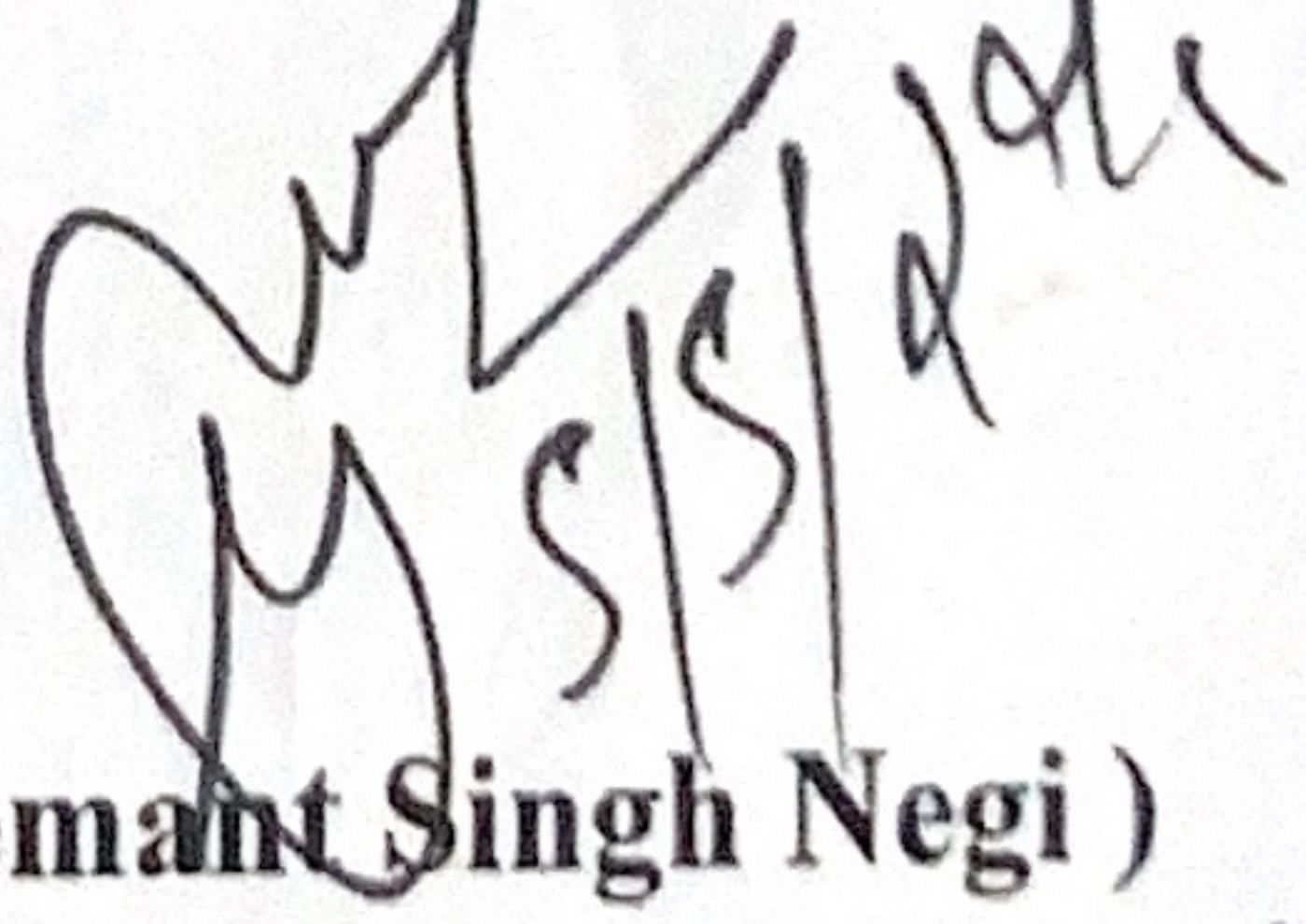
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(Mfg) Garhwal Mandal
Uttarakhand



Form -26

(See Rules 73 and 83)

**Certificate of renewal of Licence to manufacture for sale of drugs
other than those specified in schedule- X**

1- Certified that Licence No. 18/UA/2012 on form 25 & 17/UA/SC/P-2012 on form 28 granted to M/s Aryan Biotech Pvt.Ltd., for the manufacture of the following categories of drugs being drugs other than those specified in Schedule C, C(1) and X & Drugs specified in Schedule C & C(1) excluding those specified in Schedule X to the Drugs & Cosmetics Rules 1945, at the premises situated at- Plot No. 16, Sector 6B, IIE, Sidcul, Ranipur, Distt. Haridwar, Uttarakhand has been renewed from: 19-04-2017 to 18-04-2022.

2- Names of approved competent technical staff:-

a. For Manufacturing:-

(1) Mr. Vishnu Goel (Tablets Capsules & Oral Liquid)

b. For Analysis: -

(1) Mr. Satya Pal Singh (Chemical, Instrumental & Microbiology)

3-Names of drugs – As per the list enclosed.

Dated: 02-11-2017



Tajber Singh
21/11/17

(Tajber Singh)
Drug Controlling &
Licensing Authority (Mfg.)
Garhwal Mandal, Uttarakhand

(Tajber Singh)
Drug Controlling & Licensing Authority (Mfg.)
Garhwal Mandal, Uttarakhand

FORM-25

[See Rule 70]

Licence to manufacture for sale [or for distribution] of drugs other than those Specified in Schedules C, C (I) and X

Number of licence and date of issue: 18/UA/2012 date of issue: 19-04-2012

1. M/s Aryan Biotech Pvt, Ltd., is hereby licensed to manufacture the following categories of drugs being drugs other than those specified in Schedules C, C (I), and X to the Drugs and Cosmetics Rules, 1945, on the premises situated at- Plot No. 16, Sector-6B, Ranipur , Haridwar, Uttarakhand under the direction and supervision of the following (Competent Technical staff:)

(a) Competent Technical Staff (Names)

(i) Manufacturing – 1. Sh. Vishnu Geol (Tablets, Capsules & Liquid Oral)

(ii) Analysis - 1. Sh. Satya Pal (Chemical, Instrumental & Microbiologist)
2. M/s Standard Lab, Delhi.

(b) Name of Drugs – As per list enclosed.

2. The licence authorizes 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 19-04-2012 to 18-04-2017

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.

Date: 19-04-2012

Signature.....

Designation.....

S. Shandil
19-04-2012
S. S. SHANDIL
Drug Licensing Authority
Garhwal Division
Uttarakhand

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.

Any change in the expert staff named in the licence shall be forthwith reported to the Licensing Authority.

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). The licence will be deemed to extend to the cosmetics 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.

