GOVERNMENT OF Haryana
Food And Drugs Administration Haryana, Sco-94,
Sector-5, Panchkula, Panchkula, Haryana-134109,
India

FORM 28

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

File No: **HR/ML/F27/2022/00013** Licence No: **MLF282022HR000002** Issue Date: **29-NOV-2022**
Site Id:HR0004379 Old Licence No: **NA** Valid Upto: **28-NOV-2027**

1. **SCOUT LIFESCIENCE PRIVATE LIMITED** is hereby licensed to manufacture at the premises situated at the **Plot No 353, Sector 2, Industrial Growth Centre HSIIDC, Saha ,Ambala ,Ambala ,Haryana ,India ,133104** the following drugs, being drugs specified in Schedules C and C(1) [excluding those specified in Schedule X] to the Drugs Rules, 1945.

Name of drugs: See **Annexure 'A'**(Total No. of Approved Products : **7**)

2. Name of approved [Competent Technical Staff]
(i).Name(s) of staff responsible for testing

Member ID	Member Name	Role	Qualification
20225927325	Ravinder	Analytical Chemist	B.Sc

(ii). Name(s) of staff responsible for manufacturing

Member ID	Member Name	Role	Qualification
20228163471	Harbir	Manufacturing Chemist	B.Sc

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 application to licence for sale.
4.The licence,unless sooner suspended or cancelled,shall remain valid perpetually.However,the compliance with the conditions 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 as 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 (23 of 1940).

Manmohan Taneja
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Manmohan Taneja
Licensing Authority

Conditions of Licence 1. This licence shall be kept on the approved premises and shall be produced at the request of an Inspector appointed under the Drugs & Cosmetics Act 1940 (23 of 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(5). 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.

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Annexure 'A' (List of Drugs)

S.No.	Generic Name	Composition
1	**Pregabalin** Capsules **(10120220900199)**	**Pregabalin , I.P., 300.0mg**
2	**Clindamycin Hydrochloride** Capsules **(10120220900189)**	**Clindamycin Hydrochloride, I.P., 300.0mg**
3	**Tetracycline Hydrochloride** Capsules **(10120220900187)**	**Tetracycline Hydrochloride, I.P., 500.0mg**
4	**Doxycycline Hydrochloride+Lactic Acid** Capsules **(10120220900227)**	**Doxycycline Hydrochloride, I.P., 100.0mg, Lactic Acid, I.P., 5.0IU**
5	**Methylcobalamin+Cholecalciferol+ Folic Acid+Pyridoxine Hydrochloride** Capsules **(10120220900190)**	**Methylcobalamin, I.P., 1500.0µg, Cholecalciferol, I.P., 1000.0IU, Folic Acid, I.P., 1.5mg, Pyridoxine Hydrochloride , I.P., 3.0mg**
6	**Methylcobalamin+Pregabalin** Capsules **(10120220900188)**	**Methylcobalamin, I.P., 750.0µg, Pregabalin , I.P., 75.0mg**

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S.No.	Generic Name	Composition
7	**Doxycycline Hydrochloride+Lactic Acid** Capsules **(10120220900226)**	**Doxycycline Hydrochloride, I.P., 100.0mg, Lactic Acid, I.P., 60.0IU**

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Note: * is applicable for export purpose only not for domestic purpose.