



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 SCHEDULE X]

File No: **HR/ML/F27/2023/00041**
Site Id:HR0022234

Licence No: **MLF282024HR000001**
Old Licence No: **NA**

Issue Date: **02-JAN-2024**
Valid Upto: **01-JAN-2029**

1. **CUCKOS PHARMACEUTICAL PVT LTD** of **CUCKOS PHARMACEUTICAL PVT LTD** is hereby licensed to manufacture at the premises situated at the **Khatoni Number 628, Kila Number 14//13, 14//12/2, 18/1, 19/1, 8/2/2, Dattaar Road, Kheri Sampla, District Rohtak, Haryana 124501, India ,Sampla ,Rohtak ,Haryana ,India ,124501** 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 : **6**)

2. Name of approved [Competent Technical Staff]

(i).Name(s) of staff responsible for testing

Member ID	Member Name	Role	Qualification
20236367134	Akshay	Analytical Chemist	B Pharma

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

Member ID	Member Name	Role	Qualification
20237686911	Yuvaraj	Manufacturing Chemist	Msc Chemistry

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).

Date:02/01/2024

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.