FORM 21B

[*See* rule 61(2)]

[1][*Licence to sell, stock or exhibit or offer for sale, or distribute*] *by wholesale drugs specified in Schedules C and C (1)* [2][*excluding those specified in Schedule X*

1. ……is hereby [1][licensed to sell, stock or exhibit or offer for sale, or distribute] by wholesale on the premises situated at the following categories of drugs specified in Schedule. C and C (1) [2][excluding those specified in Schedule X] to the Drugs and Cosmetics Rules, 1945.

Categories of drugs

2. This licence shall be in force from..... to…….

[3][2A. The sale shall be made under the personal supervision of a competent person.
(Name of the competent person)].

3. This licence is subject to the conditions stated below and to the provisions of the Drugs and Cosmetics Act, 1940 and the rules thereunder.

Licence No ……..

Date..... *Licensing Authority.*

Conditions of Licence

1. This licence shall be displayed in a prominent place in a part of the premises open to the public.

[4][2.***]

3. If the licensee wants to sell, stock or exhibit for sale or distribute during the currency of the licence additional categories of drugs listed in Schedules C and C (1) [2][excluding those specified in Schedule X] but not included in this licence, heshould apply to the Licensing Authority for the necessary permission. This licence will be deemed to extend to the categories of drugs in respect of which such permission is given. This permission shall be endorsed on the licence by theLicensing Authority.

[5][4. (i) No drug shall be sold unless such drug is purchased under a cash or credit memo from a duly licensed dealer or a duly licensed manufacturer.

(ii) No sale of any drug shall be made for purposes of resale to a person not holding the requisite licence to sell, stock or exhibit for sale or distribute the drug:

Provided that this condition shall not apply to the sale of any drug to —

(a) an officer or authority purchasing on behalf of Government, or

(b) a hospital, medical, educational or research institute or a registered medical practitioner for the purpose of supply to his patients, or

[6][(c) a manufacturer of hydrogenated vegetable oils, beverages, confectionary and othernon-medicinal products, where such drugs are required for processing these products.]

[7][5.***]

[8][6. 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 Licensing Authority in the name of the firm with the changed constitution.

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1. Subs. by. G.S.R.788(E), dt. 10.10.1985.
2. Subs. by.G.S.R.462(E), dt. 22.6.1982
3. Ins. by.G.S.R. 681(E), dt. 6.6.1988.
4. Condition no. 2 omitted by.G.S.R. 17(E), dt. 7.1.1986
5 Added by. Notfn. No. F. 1-63/61 -D, dt. 17.7.1963.
6.Added by Notfn. No. F. 1-113/69-D, dt. 23.12.1969.
7. Condition 5 omitted by S.O. 289, dt 20.12.1973 (w.e.f. 3.2.1973)
8. Ins. By S.O. 1458, dt:27.4.1965.