Form 27-D
[See Rule 75]

Application for grant [***] of a licence to manufacture for sale or for
distribution of Large Volume Parenterals/Sera and Vaccines excluding those
specified in Schedule X

1. I/We........................................ of ........................................ hereby apply for the grant [***] of a licence to
manufacture for sale or distribution on the premises situated at
........................................ the under
mentioned Large Volume Parenterals / Sera and Vaccines, specified in
Schedule C and C(1), to the Drugs and Cosmetics Rules, 1945.

2. Name(s) of Drug(s)
........................................................................................................
................................................................
...........................................................................
(each item to be separately specified).

3. The name(s), qualification and experience of the competent technical
staff responsible
   for the manufacture of the above-mentioned drugs.
   (a) Name(s) of staff responsible for testing:

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........................................................................

   (b) Name(s) of staff responsible for manufacturing:

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........................................................................

4. The premises and plant are ready for inspection/will be ready for
inspection on ..........................................................
........................................................................................................

5. A fee of rupees ......................... has been credited to Government under the
head of account ......................

Date:......................  Signature:......................
Designation ..............

*** Omitted by GSR 1337 (E) dt. 27.10.2017
Notes:

1. The application is to be accompanied by a plan of the premises, list of equipment and machinery to be employed for manufacture and testing; memorandum of association / constitution of the firm; copies of qualification and experience of competent technical staff and documents relating to ownership or tenancy of the premises.

2. A copy of the application together with relevant enclosures shall also be sent each to Central Licence Approving Authority and concerned Zonal / Sub-Zonal Officers of the Central Drugs Standard Control organisation.