Application for grant/renewal of licence to manufacture Blood products for sale or distribution

1. I/We................................................................. of M/s................................ hereby apply for the grant of licence/renewal of licence number ............... dated ............... to manufacture products on the premises situated at .................................................................

2. Name(s) of item(s).
   1. ........................................................................
   2. ........................................................................
   3. ........................................................................
   4. ........................................................................

3. The name(s), qualification and experience of competent technical staff as under.
   (a) responsible for manufacturing
   (b) responsible for testing
      1 ........................................ 1 ...........................
      2 ........................................ 2 ...........................
      3 ........................................ 3 ...........................

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

5. A licence fee of rupees ..................................... and an inspection fee of rupees ..................................... has been credited to the Government under the head of account ............... (receipt enclosed).

Dated..............
Signature..............
Name & Designation............................................
* Delete whichever is not applicable.

Note.1: The application shall be accompanied by a plan of the premises, list of machinery and equipments for manufacture of blood products, memorandum of association / constitution of the firm, copies of certificate relating to educational qualification and experience of the competent technical staff and documents relating to ownership or tenancy of the said premises.

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