Dr. B.Anjaiah, M.D., (Paed.), D.CH,
Director.

FORM - 28-C

[See Rule 122-G]

LICENCE TO OPERATE A BLOOD BANK FOR COLLECTION, STORAGE AND PROCESSING OF WHOLE HUMAN BLOOD AND/OR ITS COMPONENTS FOR SALE OR DISTRIBUTION.

1. No. of Licence: **04/PRK/AP/2011/BB/G** Date of issue: 19 -05-2011 at the premises situated at Ongole, Prakasam District, Andhra Pradesh.

2. M/s. Rajeev Gandhi Institute of Medical Sciences Hospital Blood Bank is hereby Licenced to collect, store, process and distribute whole Blood and/or its components.

3. Name(s) of the Item(s) : WHOLE HUMAN BLOOD I.P.

4. Name(s) of Competent Technical staff :

   i) Dr. V. Aruna Kumari, M.D. (Path) : Medical Officer.
   ii) Smt. I. Anuradha : Regd. Nurse.
   iii) Smt. S. Rani : Regd. Nurse.
   iv) Smt. S. Sudha Rani : Regd. Nurse.
   v) Smt. B.V. Rama Lakshmi : Regd. Nurse.
   vi) Smt. A. Vara Lakshmamma : Regd. Nurse.
   vii) Mr. M. Bala Krishna, DMLT : Blood Bank Technician.

5. The Licence authorises Licensee to collect, store, distribute and processing of Whole Blood and/or Blood Components subject to the conditions applicable to this Licence.

6. The Licence shall be in force from 19 -05-2011 to 18 -05-2016.

7. The Licence shall be subject to the conditions stated below and to such other conditions as may be specified from time to time in the rules made under the Drugs and Cosmetics Act, 1940.

Dated: -05-2011.

Signature : 11/5/11

Name and Designation :

Licensing Authority :

JOINT DIRECTOR
DRUGS CONTROL ADMN.,
GOVT. OF A.P.
HYDERABAD - 500 038

DIRECTORATE OF DRUGS CONTROL ADMINISTRATION GOVERNMENT OF ANDHRA PRADESH HYDERABAD

// CENTRAL LICENCE APPROVING AUTHORITY //

डा. सुरिन्द्र सिंह
Dr. SURINDER SINGH
औषधि नियंत्रक (भारत)/Drugs Controller (India)
स्वास्थ्य सेवा महानिदेशालय
Dte. General of Health Services
FDA Bhawan, Kotla Road,
New Delhi-110002

Conditions over leaf.....

CONDITIONS OF LICENCE

1. The licencee shall neither collect Blood from any professional donor or paid donor nor shall be prepare blood components from the blood collected from such a donor.

2. The licence and any certificates of renewal in force shall be displayed on the approved premises and the original shall be produced at the request of an Inspector appointed under the Drugs and Cosmetics Act, 1940.

3. Any change in the Technical staff shall be forthwith reported to the Licensing Authority and/ or Central Licence Approving Authority.

4. The Licencee shall inform the Licensing Authority and/or Central Licence Approving 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 falid for maximum period of three months from the date on which the changes has taken place unless, in the meantime, a fresh licence has been taken from the licensing Authority and/or Central Licence Approving Authority in the name of the firm which changed constitution.