File No. CLAA/B&BP/TEL/06/2025-D
Government of India
Directorate General of Health Services
Central Drugs Standard Control Organisation (HQ)
(Blood Centre Division)

FDA Bhawan,
Kotla Road, New Delhi -110002
Dated:

To,

The Deputy Director, Nizamabad,
Licensing & Controlling Authority,
Drugs Control Administration,
D. No. 6-2-413, First Floor, Subhash Nagar,
Nizamabad, Telangana-State.

Subject: Approval of grant of Blood Centre license for the collection, processing, storage and distribution of Whole Human blood I.P. only issued to M/s Raja Rajeshwari Hospital Blood Centre (A Unit of Maheshwara Educational Society), Sy No. 473 Ground Floor Chitkul V Isnapur X Roads, Patancheru M Sanga Reddy District Telangana, Patancheru, District-Sangareddy, State-Telangana, India -502307–Regarding.

Reference no. – ONDLS File No.: TG/BB/F27C/2025/00051.

Sir,

With reference to the letter cited above on the subject matter, CLAA hereby approves the grant of Blood Centre license for the collection, processing, storage and distribution of Whole Human blood I.P. only issued to M/s Raja Rajeshwari Hospital Blood Centre (A Unit of Maheshwara Educational Society), Sy No. 473 Ground Floor Chitkul V Isnapur X Roads, Patancheru M Sanga Reddy District Telangana, Patancheru, District-Sangareddy, State-Telangana, India -502307–Regarding.

1. The Blood Centre may be re-inspected periodically at least once in a year from the date of licensing by a team comprising of Drugs Inspectors of CDSCO and State Licensing Authority and if required with an expert. The report may be forwarded to this Directorate for information.

2. The Blood Centre should comply with all the provisions prescribed under schedule F Part XII-B of the Drugs Rules, 1945.

Yours faithfully,

CHANDRASHEKAR RANGA

Digitally signed by CHANDRASHEKAR RANGA
Date: 2025.07.04 11:08:10 +05'30'

(Dr. Ranga Chandrashekar)
Joint Drugs Controller (India) &
Central License Approving Authority

Central Drugs Standard Control Organization
Ministry of Health and Family Welfare, Directorate General of Health Services, Government of India FDA Bhavan, ITO, Kotla Road, New Delhi - 110002,DELHI, New Delhi, Delhi-110002, India

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FORM 28C

(See rule 122-G)

Licence to operate a Blood Centre for collection, storage and processing of whole human blood and/or it's components for Sale or Distribution

File No: **TG/BB/F27C/2025/00051** Licence No: **BBF28C2025TG000062** Site Id:TG0000361
Old Licence No: NA

1. The premise situated at **SY NO 473 GROUND FLOOR CHITKUL V ISNAPUR X ROADS PATANCHERU M SANGA REDDY DISTRICT TELANGANA ,PATANCHERU ,Sangareddy ,Telangana ,India ,502307**.
2. M/s **RAJA RAJESHWARI HOSPITAL BLOOD CENTRE (A UNIT OF MAHESHWARA EDUCATIONAL SOCIETY)** is hereby licensed to collect, store, process and distribute whole blood and/or its components.
3. Name(s) of the item(s): (See Annexure 'A')
4. Name(s) of Competent Technical Staff

Member ID	Member Name	Role / Designation	Qualification
20252295965	Ms. Pavaleena Pothuraj	Technical Supervisor	B.Sc MLT
20252703156	Ms. Shoba RANI B	Registered Nurse	GNM
20253753456	Ms. Sangeetha Marriwar	Blood Centre Technician	Diploma in medical laboratory technology
20254611105	Ms. Renuka Kuppala	Blood Centre Technician	Diploma in medical laboratory technology
20257175571	Dr. Pooja PREETHAM Ummadisetti	Medical Officer	MBBS

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 License shall be in force from the date of approval of CLAA which is valid up to 5 years
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.

Condition of Licence

1.The licensee shall neither collect blood from any professional donor or paid donor nor shall he prepare blood components from the blood collected from such a donor.
2. The licence and any certificate 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 licensee 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 valid for maximum period of three months from the date on which the change 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 with the changed constitution.

10/6/2025

CHANDRASHEKAR RANGA Digitally signed by CHANDRASHEKAR RANGA Date: 2025.07.04 11:08:30 +05'30'

Licensing Authority (Telangana)

Central Licence Approving Authority (CDSCO)

Date: 10-Jun-25

B. SARALA B.Pharm
Deputy Director, Nizamabad
Licensing & Controlling Authority
Drugs Control Administration
Hyderabad, Telangana-State

Central Drugs Standard Control Organization
Ministry of Health and Family Welfare, Directorate General of Health Services, Government of India FDA Bhavan, ITO, Kotla Road, New Delhi - 110002,DELHI, New Delhi, Delhi-110002, India

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Annexure 'A'(Name of Items)

S.No	Item Name:
1	WHOLE HUMAN BLOOD I.P , I.P.

10/6/2025

Licensing Authority
(Telangana)

CHANDRASHEKAR RANGA
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Central Licence Approving Authority
(CDSCO)

Date: 10-Jun-25

P. SARALA B.Pha
Deputy Director, Nizamaba
Licensing & Controlling Authority
Drugs Control Administratio
Hyderabad, Telangana-S

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