Certificate of Compliance

CE

We hereby declare that the technical file of product complied with the requirement of directives Pressure Equipment Directive (PED) 2014/68/EU And Machinery directive- 2006/42/EC.

Certificate No.: CE-5213

Name : EL SHADDAI REFRIGERATION & AIR CONDITIONING PVT. LTD.

Address : NO.2/991, 3RD FLOOR, KESAVA MANSION, 2ND CROSS, 3RD MAIN, 1ST BLOCK, 4TH STAGE, BTM LAYOUT, BANGALORE, KARNATAKA - 560076, INDIA

Product : Design, Development, Manufacturing, Exporting Installation And Servicing Of Medical Equipment, Hospital Equipment, Refrigeration & Airconditioning products as receiver, Accumulators, Evaporator, Compressor Rack, Condensing Unit, Water-Cooled Condenser, Air-Cooled Condenser, Incubator, Oil Separator, Oil Receiver, Helical Oil Separator, Process Chiller, Heat Pump, Control Panel, Solar/Wind DC Driven Cold Room, Ice Lined Refrigerator, Active Cold Box/Cold Box, Walk In Cooler/Cold Room, Walk In Freezer, Deep Freezer, Ultra Low Temperature Deep Freezer, Blood Storage Refrigerator, Bio Safety Cabinet, Laminar Air Flow, Platelet Incubator And Agitator, Blood Donor Chair, Shock Freezer/Blast Freezer, Blood Collection Monitor, Refrigerated Centrifuge, Blood Tube Sealer, Bod Incubator, CO2 Incubator, Vaccine Carrier, and Laboratory Refrigerator

Complies with the requirements applicable to it

The Certification body has performed an audit of the above products quality system covering the design, manufacture and final inspection of the certified products. The quality system has been assessed, approved and is subject to continuous surveillance according to the directives Pressure Equipment Directive (PED) 2014/68/EU And Machinery directive- 2006/42/EC.

This certificate is issued under the following conditions:

1. It applies only to the quality system maintained in the manufacture of above referenced models and it does not substitute the design or type-examination procedures, if requested.
2. The certificate remains valid until the manufacturing conditions or the quality systems are changed.
3. The certificate validity is conditioned by positive results or surveillance audits.
4. The CE mark as shown above can be used, under the responsibility of the manufacturer, after completion of an EC Declaration of conformity and compliance with all relevant EC Directives. The statement is based on a single evaluation of one sample of above mentioned product. It does not imply an assessment of the whole production.

Validity of this certificate can be verified at www.rbscert.com/verify

Date of initial registration	28th JAN 2025
1st Surveillance Audit due	27th JAN 2026
2nd Surveillance Audit due	27th JAN 2027
Recertification due	27th JAN 2028

(Subject to the company maintaining its system to the required standard)

Validity of this certificate is subject to Annual Surveillance audits to be done successfully on or before 365 days from date of the Audit. (In case Surveillance Audit is not allowed to be conducted; this certificate shall be Suspended/ withdrawn).

The validity of this certificate can be verified at www.rbscert.com

Email :- info@rbscert.com website : www.rbscert.com

RBS Quality Certification Pvt. Ltd. is Accredited by IAF-UK Ltd. (www.iaf-uk.co.uk)
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Director