



Certificate of Compliance

MANUFACTURER

Name : M/S RESPROEX 2

Address : C-916 2ND FLOOR, SECTOR 7, DWARKA, NEW DELHI - 110077

The manufacturer's technical documentation as required for Medical Council according to the directive 93/42/EEC has been reviewed and found to comply with the requirements for Council Directive on Medical Devices 93/42/EEC as Amended 2007/47/EC. Any significant changes in the design or construction of the product, not agreed upon by us, this declaration will lose its validity.

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 not changed.

The CE mark as shown above can be used, under the responsibility of the manufacturer, 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.

Certificate No : UKAAB/DXPUL3082J

Issued Date : 20/09/2021

Expiry Date : 19/09/2024

1st Surveillance : 19/09/2022

2nd Surveillance : 19/09/2023



Authorised Signatory



Accredited by:

UK Akkreditation and Certification Board Limited

Address: 2nd Floor, College House, 17 King Edwards Road, Ruislip, London, United Kingdom, HA4 7AE (UK)

Registered with the Registrar of Companies for England and Wales (UK), Registration No. 12773650, www.ukaab.org.uk

MLR Certification Private Limited ,

Corporate Address: C-2/170, 3rd Floor, New Ashok Nagar, New Delhi-110096, INDIA

Regd. Address: 1/11, Ground Floor, Virat Khand-1, Gomti Nagar, Lucknow, Uttar Pradesh-226010, INDIA

Registered with Ministry of Corporate Affairs, Government of India, CIN-U74999UP2020PTC136667

This certificate is the intellectual property of MLR Certification Private Limited and can be maintained through surveillance and renewal audits, and shall be returned immediately upon request. The approval is subject to the company maintaining its system to the required standard, which will be monitored by MLR Certification Private Limited. This certificate includes declaration of manufacturer/firm/company that they are following the applicable guidelines/standards.

Validity of this certificate can be verified at www.mlrcertification.com





Certificate of Compliance

ANNEXURE-1

Certificate No : UKAAB/DXPUL3082J

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PRODUCT GROUP :

TRANSPORT VENTILATOR, DEFIBRILLATOR, AUTOMATIC EXTERNAL DEFIBRILLATOR, SYRINGE INFUSION PUMP, VOLUMETRIC INFUSION PUMP, CARDIAC PATIENT MONITOR, PATIENT MONITOR, MONITOR, CAPNOMETER, OXIMETER, PULSE OXIMETER, FINGER TIP OXIMETER, PATIENT HANDLING SYSTEM AND AMBULANCE ACCESSORIES, ELECTRIC AND MANUAL SUCTION MACHINES AND ASPIRATORS

MODELS :

R-VAC, REX-VAC, SUPREX, SUPREX PLUS, REX12, REX 8, REX-PRO, REXPRO-EX, REVIVIFYING, REINVIGORATE, RESTORATIVE, INVIGORATING, IMPROVE- CRISIS, AMELIORATE- CATASTROPHE, LAZARETTO, LAZARETTO-PRO, LAZARETTO-PLUS, LAZARETTO-EXPERT, PARAMEDICS-ONE, PARAMEDICS-PRO, PARAMEDICS-PLUS, PARAMEDICS-EX, PARAMEDICS-II, PULSE-EX, PULSE-PRO, HEART-EX, HEART-PLUS, HEART-PRO, HEART-XL, REX-JET



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