



Certificate of Compliance

This is to Certify that
The Quality Management System for Medical Devices of

M/S RESPROEX 2

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

has been independently assessed and found to be in compliance
with the requirement of the standard

ISO 13485:2016

For the following Scope :

Design, Development, Manufacturing and Trading of Ambulances, Ambulance Components, Ambulance Stretchers, Specialty Vehicles, Mobile Medical Units, Ambulance Medical Equipment's and Devices, Hospital Emergency and Rescue Equipment's

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

Certificate No : UKAAB/CYOTG9615L

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

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

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