

Certificate of Registration

Compliance

CE

We hereby declare that the technical file of product complied with the requirement of directives 2014/35/EU Low Voltage Directive

COMPANY NAME - AAR TECH MEDICAL SYSTEM
REGD. - H-237, HALDONI, KULESHRA, GREATER NOIDA, GAUTAM
BUDDHA NAGAR-201305, UTTAR PRADESH, INDIA

THIS CERTIFICATE REFERRED TO ABOVE COVERS THE FOLLOWING PRODUCTS LIKE:

Product Group:- MEDICAL & SURGICAL PRODUCTS

- Wall Paneling
- Ceiling Paneling
- Pressure Relief Dampers
- Laminar Air Flow & Plenum Systems
- Surgeon Control Panel - Membrane & Touch Screen
- Automatic Hermetically Sealing Sliding Doors Hospital Doors- Flap, Automatic & Manual
- Pressure Relief Dampers
- X-Ray Viewing Screen
- Single Arm & Double Arm Pendants
- Surgical Scrub Sink
- High Definition Recorder
- Corner Guard/Crash Guard Rail

The Certification body has performed an audit of the above product Quality system covering the design, manufacture and final inspection of the certified product. The Quality system has been assessed, approved and is subject to continuous surveillance according to The Low Voltage Directive (2014/35/EU)

Thomas disuja

Authorized Signatory

British of Columbia Inspection & Certification



📍 Head office Address : 7445 132 St Suite 1001, Surrey, BC V3W 1J8, Canada

COMPANY NAME - AAR TECH MEDICAL SYSTEM
REGD. - H-237, HALDONI, KULESHRA, GREATER NOIDA,
GAUTAM BUDDHA NAGAR-201305, UTTAR PRADESH,INDIA

THIS CERTIFICATE REFERRED TO ABOVE COVERS THE FOLLOWING PRODUCTS LIKE:

Product Group:- MEDICAL & SURGICAL PRODUCTS

- Fully & Semi-Automatic Control Panel-Oxygen System / Nitrous-Oxide System /Carbon Dio-Oxide System
- Manifold System - Oxygen / Carbon Dio-Oxide
- Emergency System - Oxygen / Carbon Dio-Oxide
- Oxygen Flow Meter with Humidifier Bottle
- Medical Gas Outlets With Probe
- Ward Vacuum Unit
- Theatre Suction Unit
- Medical Gas Alarm (Main & Area)
- Valve Boxes
- Bed Head Panels - Horizontal & Vertical
- High Pressure Tubes
- Curtain Track
- IV Tree

The Certification body has performed an audit of the above product Quality system covering the design, manufacture and final inspection of the certified product. The Quality system has been assessed, approved and is subject to continuous surveillance according to The Low Voltage Directive (2014/35/EU)

The 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. After fulfilling the relevant Standard testing performance, the manufacturer shall affix to each device, of the referenced models

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 <https://www.bccert.org/searchresult.php>

Certificate issue Date : 08/05/2024 | Certificate Expiry Date : 08/04/2027
1st Surveillance due Before : 08/05/2025 | 2nd Surveillance due Before : 08/05/2026
Certificate Number: BCI-15013CE

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