



Certificate of Registration

This is to certify that

MEDTECH DEVICES

F-45, 2ND FLOOR, OKHLA INDUSTRIAL AREA, PHASE-1, NEW DELHI-110020, INDIA
has been independently assessed by QRO
and is compliant with the requirement of:

ISO 9001:2015

Quality Management System

For the following scope of activities:

MANUFACTURER, SUPPLIER & EXPORTER OF MEDICAL DEVICES (HEMODIALYSIS CATHETER KIT & ACCESSORIES - LONG TERM & SHORT TERM, CVC KITS & ACCESSORIES, MANIFOLD & ACCESSORIES, GUIDEWIRES, PTCA KIT & ACCESSORIES, ANGIO KIT & ACCESSORIES, INTRODUCER SHEATH KIT & ACCESSORIES, BLOOD LINE SET, A.V. FISTULA NEEDLE, INTRODUCER NEEDLE, C.T. CONNECTION TUBE (PLAIN & SPIRAL), EXTENSION LINE, HIGH PRESSURE TUBING, PRESSURE MONITORING LINE, HIGH PRESSURE SYRINGES, SPIKE, J TUBE FOR C.T. & ANGIO, BIOPSY GUN & BIOPSY NEEDLE, PROCEDURE KIT / DRESSING KIT, CONTROL SYRINGES, TRANSDUCER PROTECTOR, INFLATION DEVICE, BONE MARROW BIOPSY SYSTEM, I.V. CANNULA, THREE WAY STOPCOCK, FOLLEY CATHETER, ORTHOPAEDIC SUCTION SET, ABSORBENT HEMOSTATIC PRESSURE DRESSING, PACING LEADS, HAND CONTROL PENCIL, DIALYZER REPROCESSOR MACHINE & ENDOTOXIN TEST MACHINE), DISPOSABLE HEMOPERFUSION CARTRIDGE, HEMOPERFUSION MACHINE, PORE CLEAN, CITRA CLEAN & UROLOGY PRODUCTS, LIQUID BANDAGE.

Date of Certification: 29th July 2025
1st Surveillance Audit Due: 28th July 2026

2nd Surveillance Audit Due: 28th July 2027
Certificate Expiry: 28th July 2028

Certificate Number: 305025072956Q




Head of Certification

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(In case surveillance audit is not allowed to be conducted; this certificate shall be suspended / withdrawn).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.qrocet.org

This certificate of registration remains the property of QRO Certification LLP, and shall be returned immediately upon request.

India Office : QRO Certification LLP

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