Certification & Inspection

Certificate of Registration

This is to certify that the
Quality Management System for Medical Devices
of

MEDTECH DEVICES

at

**F-45, 2ND FLOOR, OKHLA INDUSTRIAL AREA, PHASE-1,
NEW DELHI-110020, INDIA**

has been independently assessed and is
compliant with the requirements of:

ISO 13485:2016

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, DIALYZER REPROCESSER MACHINE & ENDOTOXIN TEST MACHINE, HAND CONTROL PENCIL, HEMOPERFUSION MACHINE, HEMODIALYSIS MACHINE, PACING LEAD, CRRT MACHINE)

Certificate Number: UQ - 2024082014

Validity of this certificate can be verified at www.ukcertifications.org.uk/verify

Date of Certification	16th August 2024
1st Surveillance Audit Due	15th August 2025
2nd Surveillance Audit Due	15th August 2026
Certificate Expiry	15th August 2027

Authorised Signatory

UKAF

UK Certification & Inspection Ltd.