# Certificate of Compliance

**Certificate Number: UQ** - 2024061722

This is to certify that

## MEDTECH DEVICES

at

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

Has successfully implemented the Quality management System and been found working satisfactorily as per the norms of **"Good Manufacturing Practice"** as laid down by "**World Health Organisation"** which has been in conformance to the requirements of

# WHO-GMP

**MANUFACTURING, EXPORT, IMPORT, MARKETING AND SALES OF MEDICAL DEVICE PRODUCTS LIKE HEMODIALYSIS CATHETERS, GUIDE WIRE, CENTRAL VENOUS CATHETER, INTRODUCER SHEATH, BIOPSYS GUN & NEEDLES, HYPODERMIC NEEDLES, MANIFOLDS, AV FISTULA NEEDLS, BLOOD TUBING SET, CT CONNECTION TUBES, MEDICAL KITS, REPROCESSER, HEMOPERFUSION AND CRRT MACHINES & OTHER MEDICAL DEVICE PRODUCTS.**

**This certificate is issued under the following conditions:**

1. It applies only to the quality system maintained in the manufacture of above referenced Models Products.
2. The certificate remains valid until the manufacturing conditions or the quality systems are changed and is subject to continuous surveillance according to the WHO-GMP Guidelines
3. The certificate validity is conditioned by positive results or surveillance audits.

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

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| Date of Certification | 17th June 2024 |
| 1st Surveillance Audit Due | 16th June 2025 |
| 2nd Surveillance Audit Due | 16th June 2026 |
| Certificate Expiry | 16th June 2027 |

Authorised Signatory