Certificate of Compliance

CE

We hereby declare that the technical file of product complied with the requirement of directive Council Directive on Medical Device Regulation -(Council Regulation 2017/745)

Certificate No.: CE-4577

Manufacturer

Name : **AHEESHA MEDICAL TECHNOLOGY PRIVATE LIMITED**

Address : **GROUND FLOOR, PLOT NO 135, PKT-H, SEC-3,BAWANA DSIDC, BAWANA, NEW DELHI, 110039, INDIA**

Product Name : **PATIENT MONITER, VESSEL SEALER, CAUTERY MACHINE, SYRINGE PUMP, INFUSION PUMP, OT TABLE, OT LIGHT, HORIZONTAL AUTOCLAVE MACHINE, VERTICAL AUTOCLAVE MACHINE, ETO MACHINE.**

Complies with the requirements applicable to it

The quality system file has been assessed, approved and is subject to continuous surveillance according to the Medical Device Regulation -(Council Regulation 2017/745)

This 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.

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 www.ukcertifications.org.uk/verify

Date of Certification	16th November 2023
1st Surveillance Audit Due	15th November 2024
2nd Surveillance Audit Due	15th November 2025
Certificate Expiry	15th November 2026

Authorised Signatory

UK Certification & Inspection Ltd.