Certificate of Compliance

CE

We hereby declare that the technical file of product complied with the requirements of EC directive 93/42/EEC Medical Device Directive (MDD).

Certificate No.: CE- 16241

Manufacturer

Name : **MANARTI EXPORTS**

Address : **Corp. Office: First Floor, Office Space No. 114-115, Vishwadeep Tower, Plot No. 4, District Center, Janak Puri, New Delhi - 110058, India**
Factory : Plot No. 78A Near Subh Nursing Home, Village Hastsal, Uttam Nagar, New Delhi - 110059, India

Products

Name : **As Per Appendix Page 1 to 12**

Complies with the requirements applicable to it

The quality system file has been assessed, approved and is subject to continuous surveillance according to the Council Directive on Medical Devices 93/42/EEC as Amended 2007/47/EC class I.

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 not changed.
3. The certificate validity is conditioned by positive results of surveillance audits.
4. 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.co.uk/verify

Date of Certification	13th July 2021
1st Surveillance Audit Due	12th July 2022
2nd Surveillance Audit Due	12th July 2023
Certificate Expiry (subject to the company maintaining its system to the required standard)	12th July 2024

Authorised Signatory

UK Certification & Inspection Ltd.