Certificate of Compliance

CE

We hereby declare that the technical file of product class 1 complied with the requirement of the Medical Devices Directives 93/42/eec of June 1993 and 89/686/eec

Applicant Certificate No: CE-9679

Name	:	**NU MED TECHNOLOGIES**
Address	:	A-227, Okhla Industrial Area, Okhla Phase-1, New Delhi-110020 (India)
Product Name	:	Manufacturing, Marketing, Development and Sales of Products for Pre-natal Care Product Categories: Fetal Heart Rate Doppler's, Fetal Monitors, Maternal & Fetal Monitors with Wired and Wireless Transducer and CTG Guided TENS Therapy for Labor Pain Management System
Brand Name "	:	"numed"
Model	:	T2FMM12 - Fetal & Maternal Monitor, T1FM12 - Fetal Monitor T1FM08 - Fetal Monitor, T1FD01 – Fetal Doppler

The Certification body has performed an audit of the above product quality system covering the design, manufacture and final inspection of the certified product. The quality system has been assessed, approved and is subject to continuous surveillance according to Directive 93/42/ec of June 1993 Medical Council Directives and 89/686/eec

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 thedesign or type-examination procedures, if requested.
2. The certificate remains valid until the manufacturing conditions orthe quality systems are changed.
3. The certificate validity is conditioned by positive results or surveillance audits.

The CE mark as shown above canbe used, under the responsibility of the manufacturer, after completion of an EC Declaration of conformity and compliance with all relevant EC Directices. The statement is based on a single evalution of one sample each of above mentioned product. It does not imply an assessment of the whole production.

Date of Certification 08th June 2021

Certificate Expiry (subject to the company maintaining its system to the required standard) 07th June 2024

Authorised Signatory

BRILLIANT CERTIFICATION & INSPECTION

Validity of this Certification is successful Annual Surveillance audits to be done.

info@bcicert.co.uk www.bcicert.co.uk

CERTIFICATE

Certificate of Registration

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

NU MED TECHNOLOGIES

at

A-227, Okhla Industrial Area, Okhla Phase-1, New Delhi
Delhi-110020 (India)

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

ISO 13485:2016

Quality Management System For Medical Devices

For the following scope of activities:

Manufacturing and Trading of Surgical Instrument, Medical Disposables, Medical Devices & Equipments

Certificate Number: UQ-482133824

Date of Certification	09th August 2020
1st Surveillance Audit Due	08th August 2021
2nd Surveillance Audit Due	08th August 2022
Certificate Expiry (subject to the company maintaining its system to the required standard)	08th August 2023

Authorised Signatory

Validity of this Certification is successful Annual Surveillance audits to be done.
This certificate is the property of Brilliant Certification & Inspection and shall be returned immediately on request, Brilliant Certification & Inspection is Accreditation by UKAF (www.ukaf.org.uk)
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