EC Certificate Full Quality Assurance System: Certificate GB19/964564

The management system of

Moor Instruments Ltd

Millwey, Axminster, Devon, EX13 5HU, UK

has been assessed and certified as meeting the requirements of

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Directive 93/42/EEC

on medical devices, Annex II (excluding Section 4)

For the following products

Laser Doppler Perfusion and Temperature Monitors and non-sterile probes
Tissue Oxygen and Temperature Monitors and non-sterile probes
Skin Heater and Temperature Monitor and non-sterile probes
Vascular Assessment Pressure Cuff Controller
Laser Perfusion Imagers
Burn Assessment Systems
Iontophoresis Controllers and non-sterile chambers
PC Software for imaging and monitoring devices.

Where the above scope includes class III medical device(s), a valid EC Design Examination Certificate according to Annex II (Section 4) is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 14 January 2020 until 15 December 2023
and remains valid subject to satisfactory surveillance audits.
Issue 2. Certified since 12 June 1998
and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered GB/PC 07523

Authorised by

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