



The management system of

Moor Instruments Ltd

Millwey, Axminster, Devon, EX13 5HU, UK

has been assessed and certified as meeting the requirements of

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

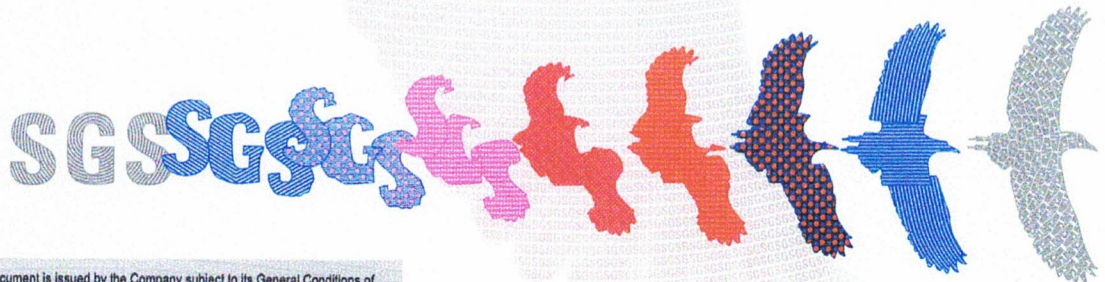
Authorised by

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LPMD5007 - Certificate CE1639 Annex II-4_EN rev. 02

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