

The management system of

MEC Medical Limited

Unit 3, Trust Estate, Wilbury Way, Hitchin, Hertfordshire, SG4 0UZ, 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

Medical Gas System Components:

Hose assemblies and Associated Fittings **Purair AGSS Systems** Medical gas Terminal units Gas specific Schrader units Ambulance change over valves Medical Gas regulators **Medical Gas Flowmeters**

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 18 May 2021 until 14 June 2023 and remains valid subject to satisfactory surveillance audits. Issue 2. Certified since 14 June 2005

> > Certification is based on reports numbered GB/PC 210478

Authorised by

Global Medical Devices Head of Notified Body

SGS Belgium NV, Notified Body 1639

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



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