EC Certificate Full Quality Assurance System: Certificate GB19/964711

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

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