Medical Device Full Quality Assurance System Certificate GB23/00000066

The management system of

MEC MEDICAL LTD

MEC House 8 Fountain Drive Hertford Hertfordshire SG13 7UB United Kingdom

has been assessed and certified as meeting the requirements of

Part II of The Medical Devices Regulations 2002, Annex II excluding section 4 [as modified by Part 2 of Schedule 2A to The Medical Devices Regulations 2002]

For the following products
Medical Gas System Components:
Hose Assemblies and Associated Fittings
Purair AGSS Systems
Medical Gas Terminal Units
Gas Specific Schrader Units
Change Over Valves
Medical Gas Regulators
Medical Gas Flowmeters

Where the above scope includes class III medical device(s), a valid Design Examination Certificate according to Annex II (Section 4) [as modified by Part 2 of Schedule 2A of The MDR 2002] is a mandatory requirement for each device in addition to this certificate to place that device on the market

Certification is based on reports numbered GB/PC/210478

Previous certificate number: N/A

Change in between this certificate and previous one: N/A

This certificate is valid from 07 June 2023 until 14 June 2028 and remains valid subject to satisfactory surveillance audits.
Issue 2. Certified since 06 February 2023

Authorised by
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Head of Approved Body 0120

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