



Borellert durch/Designated by
 Zertifikat der Länder
 für Gesundheitsschutz
 bei Eingriffen in
 Medizinprodukte
 ZLG-B5-144, 10.0



Product Service

EC Certificate

Full Quality Assurance System
 Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
 (Devices in Class IIa, IIb or III)

No. G1 010578 0037 Rev. 01

Manufacturer:

Drägerwerk AG & Co. KGaA
 Moislinger Allee 53-55
 23542 Lübeck
 GERMANY

Facility(ies):

Drägerwerk AG & Co. KGaA
 Revalstraße 1, 23560 Lübeck, GERMANY

Drägerwerk AG & Co. KGaA
 Moislinger Allee 53-55, 23542 Lübeck, GERMANY

Product Category(ies):

Anaesthetic equipment with standard accessories,
 Infusion equipment with standard accessories,
 Pediatric equipment with standard accessories,
 Lung ventilator equipment with standard accessories,
 Monitoring equipment with standard accessories,
 Equipment for suction, breathing-, inhalation-, oxygen-
 and aerosol-therapy with standard accessories,
 Medical supply units and terminal units for pressurized
 medical gases and vacuum,
 Pipelines for compressed medical gases and vacuum,
 Anaesthetic gas scavenging systems, Components for
 medical gas management systems, Software for diagnosis based on clinical
 data incl. patient data, monitoring and device parameter, Visualization,
 diagnostic and therapeutic software for anaesthesia and respiratory devices

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned
 manufacturer has implemented a quality assurance system for design, manufacture and final
 inspection of the respective devices / device categories in accordance with MDD Annex II. This quality
 assurance system conforms to the requirements of this Directive and is subject to periodical
 surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See
 also notes overleaf.

Report No.:

713182398

Valid from:

2020-01-15

Valid until:

2024-05-26

Date,

2019-12-09

Christoph Dicks
 Head of Certification/Notified Body