

Drägerwerk AG & Co. KGaA, 23542 Lübeck

To whom it may concern

#### Manufacturer's Authorization

January 27, 2021

We, Drägerwerk AG & Co. KGaA, Moislinger Allee 53-55, 23558 Lübeck, Germany, who is an established and reputable manufacturer of medical equipment, having factories at Lübeck (Germany), Telford (United States), Andover (United States) and Shanghai (China), do hereby declare that

> "Echipamed-Plus" SRL Valea Trandafirilor 24 "B", of, 80 MD-2001, Chisinau Republic of Moldova

is our Independent Re-Seller for Anesthesia Devices, Incubators, Ventilation Machines, Monitoring, Pendant and Operating Lights of Drägerwerk AG & Co. KGaA in the territory of the Republic of Moldova.

We declare that the above mentioned company is authorized to quote, sell, subsequently negotiate and sign contracts, as well as to perform installation and after sales service of Anesthesia Devices, Incubators, Ventilation Machines, Monitoring, Pendant and Operating Lights manufactured by us in their own name and on their own account.

This authorization letter will remain valid until 31.12.2021.

Duly authorized to sign this Authorization on behalf of

Drägerwerk AG & Co. KGaA

Thomas Engler

CEO Region Europe

Martin Koch

MD Direct Export

Drägerwerk AG & Co. KGaA Moislinger Allee 53-55 23558 Lübeck, Germany Postal address: 23542 Lübeck, Germany Tel +49 451 882-0 Fax +49 451 882-2080 info@draeger.com www.draeger.com VAT no. DE135082211

Bank details: Commerzbank AG, Lübeck IBAN: DE95 2304 0022 0014 6795 00 Swift-Code: COBA DE FF 230 Sparkasse zu Lübeck IBAN: DE15 2305 0101 0001 0711 17 Swift-Code: NOLADE21SPL

Registered office: Lübeck Commercial register: Local court Lübeck HRB 7903 HL General partner: Drägerwerk Ven

Registered office: Lübeck Commercial register: Local court Lübeck HRB 7395 for Drägerwerk AG & Co. KGaA and Dragerwerk Verwaltungs AG:

Chairman of the Supervisory Board



# **CERTIFICATE**

The Certification Body of TÜV SÜD Management Service GmbH

certifies that

## Dräger

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

for the Scope of application

Design and development, production and distribution of diagnostic and therapeutic medical devices and installations as well as consulting and services in the field of medical technology

Revalstraße 1, 23560 Lübeck Germany

for the Scope of application

Production and distribution of diagnostic and therapeutic medical devices and installations

has established and applies a Quality Management System.

An audit was performed, Order No. 707037695.

Proof has been furnished that the requirements according to

ISO 9001:2015

are fulfilled.

The certificate is valid from 2021-01-15 until 2024-01-14.
Certificate Registration No.: 12 100 49423 TMS.

Head of Certification Body Munich, 2021-01-13









#### Certificate

No. Q5 010578 0031 Rev. 01

Holder of Certificate: Drägerwerk AG & Co. KGaA

Moislinger Allee 53-55 23542 Lübeck

GERMANY

**Certification Mark:** 



Scope of Certificate: Design, Development, Manufacture and

Distribution of Diagnostic and Therapeutic Medical Devices and Installations as well as

Consulting and Services in the Field of

Medical Technology. Diagnostic and Therapeutic Medical Devices and

Installations: Anaesthetic Equipment,
Infusion Equipment, Pediatric Equipment

for Warming- and Photo-Therapy, Lung

Ventilator Equipment, Monitoring

Equipment, Clinical Decision Support Software, Patient Data Management

Software, Equipment for Suction,

Breathing-, Inhalation-, O2- and Aerosol-Therapy, Medical Gas Management and

Supply Systems as well as Medical Lights

The Certification Body of TÜV SÜD Product Service GmbH certifies that the company mentioned above has established and is maintaining a quality management system, which meets the requirements of the listed standard(s). All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:Q5 010578 0031 Rev. 01

Report No.: Valid from: 713193628 2021-01-18

Valid until: 2024-01-13

Christoph Dicks

Head of Certification/Notified Book

Date,

2021-01-18

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany



#### Certificate

No. Q5 010578 0031 Rev. 01

Applied Standard(s): EN ISO 13485:2016

Medical devices - Quality management systems -

Requirements for regulatory purposes

(ISO 13485:2016) DIN EN ISO 13485:2016

Facility(ies):

Drägerwerk AG & Co. KGaA

Moislinger Allee 53-55, 23542 Lübeck, GERMANY

Design, Development, Manufacture and Distribution of Diagnostic and Therapeutic Medical Devices and Installations as well as Consulting and Services in the Field of Medical Technology. Diagnostic and Therapeutic Medical Devices and Installations: Anaesthetic Equipment, Infusion Equipment, Pediatric Equipment for Warming- and Photo-Therapy, Lung Ventilator Equipment, Monitoring Equipment, Clinical Decision Support Software, Patient Data Management Software, Equipment for Suction, Breathing-, Inhalation-, O2- and Aerosol-Therapy, Medical Gas Management and Supply Systems as well as Medical Lights

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

Manufacture and Distribution of Diagnostic and Therapeutic Medical Devices and Installations as well as Consulting and Services in the Field of Medical Technology. Diagnostic and Therapeutic Medical Devices and Installations: Anaesthetic Equipment, Infusion Equipment, Pediatric Equipment for Warming- and Photo-Therapy, Lung Ventilator Equipment, Monitoring Equipment, Clinical Decision Support Software, Patient Data Management Software, Equipment for Suction, Breathing-, Inhalation-, O2- and Aerosol-Therapy, Medical Gas Management and Supply Systems as well as Medical Lights

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### CERTIFICATE

Management system as per

ISO 14001: 2015

In accordance with TÜV NORD CERT procedures, it is hereby certified that

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

with the site Revalstraße 1, 23560 Lübeck, Germany

applies a management system in line with the above standard for the following scope

Design and development, manufacturing, sales and servicing of diagnostic and therapeutic medical devices and installations as well as consulting and services in the field of medical technology

Certificate Registration No. 07 104 980284-050 Audit Report No. 3522 3922 Valid from 2018-12-16 Valid until 2021-12-15 Initial certification 2009-12-16

Certification Body at TÜV NORD CERT GmbH Essen, 2018-11-08

This certification was conducted in accordance with the TÜV NORD CERT auditing and certification procedures and is subject to regular surveillance audits. This certificate is valid in conjunction with the main certificate.

TÜV NORD CERT GmbH

Langemarckstraße 20

45141 Essen











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-, oxygenand 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 scaving 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 anesthesia 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.:

713162398

Valid from:

2020-01-15

Valid until:

2024-05-26

Date.

2019-12-09

Christoph Dicks

Head of Certification/Notified Body

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TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339