

Drägerwerk AG & Co. KGaA, 23542 Lübeck

To whom it may concern

Our reference

DW-legal / ew-ro / 119/13

Phone

+49 451 882-2471

Fax

+49 451 882-72471

E-mail

Erika.Wagner@draeger.com

February 27, 2019

## Manufacturer's Authorization

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, 24B, of. 80**  
**MD-2001 Chisinau**  
**Republic of Moldova**


is an official distributor for Anesthesia Devices, Incubators, Ventilation Machines, Monitoring, Pendant and Operating Lights of Drägerwerk AG & Co. KGaA in the territory of the Republic of Moldova.

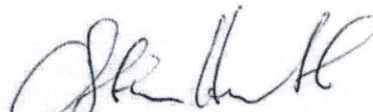
We declare that above mentioned company is authorized to quote, sell, subsequently negotiate and sign contracts, as well as to perform installation and after sales service of Anesthesia Devices, Incubators, Ventilation Machines, Monitoring, Pendant and Operating Lights manufactured by us in their own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

This authorization letter will remain valid until 31.12.2019.

Duly authorized to sign this Authorization on behalf of:

**Drägerwerk AG & Co. KGaA**

  
Dr. Erika Wagner  
authorized representative

  
Dr. Christian Hauswaldt  
authorized representative



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 33 03  
Sparkasse zu Lübeck  
IBAN: DE15 2305 0101 0001 0711 47  
Swift-Code: NOLADE21SR1

Registered office: Lübeck  
Commercial register:  
Local court Lübeck HRB 7903 HL  
General partner: Drägerwerk Verwaltungs AG  
Registered office: Lübeck  
Commercial register:  
Local court Lübeck HRB 7395 HL

Chairman of the Supervisory Board  
for Drägerwerk AG & Co. KGaA  
and Drägerwerk Verwaltungs AG:  
Stefan Lauer  
Executive Board:  
Stefan Dräger (chairman)  
Rainer Klug  
Gert-Hartwig Lescow  
Dr. Reiner Piske  
Anton Schrofner







Management Service

# CERTIFICATE

The Certification Body  
 of TÜV SÜD Management Service GmbH  
 certifies that

**Dräger**

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

has established and applies  
 a Quality Management System for

**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.**

An audit was performed, Report No. **707037695**.

Proof has been furnished that the requirements  
 according to

**ISO 9001:2015**

are fulfilled.

The certificate is valid in conjunction  
 with the main certificate from **2018-01-15** until **2021-01-14**.

Certificate Registration No.: **12 100 49423/01 TMS**.



*M. Wege*

Product Compliance Management  
 Munich, 2018-01-09







Product Service

# CERTIFICATE

No. Q5 17 11 10578 031

Holder of Certificate: **Drägerwerk AG & Co. KGaA**
**Dräger**

 Moislinger Allee 53-55  
 23542 Lübeck  
 GERMANY

**Facility(ies):**

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

 Drägerwerk AG & Co. KGaA  
 Revalstraße 1, 23560 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

**Applied Standard(s):**

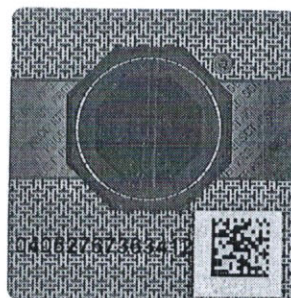
 EN ISO 13485:2016  
 Medical devices - Quality management systems - Requirements for regulatory purposes  
 (ISO 13485:2016)  
 DIN EN ISO 13485:2016

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). See also notes overleaf.

 Report No.: 713113147  
 Valid from: 2018-01-14  
 Valid until: 2021-01-13

Date, 2017-12-27

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DAKKS

 Deutsche  
 Akkreditierungsstelle  
 D-ZM-11321-01-00

TÜV SÜD Product Service GmbH · Zertifizierungsstelle · Ridlerstraße 65 · 80339 München · Germany

TÜV®





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 14 12 10578 004

**Manufacturer:****Drägerwerk AG & Co. KGaA**

Moislinger Allee 53-55  
23542 Lübeck  
GERMANY

**Facility(ies):**

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

Drägerwerk AG & Co. KGaA  
Revalstraße 1, 23560 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

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.:**

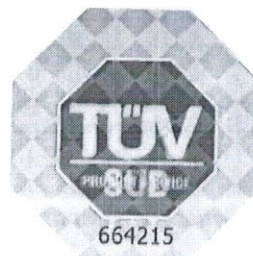
713052642

**Valid from:**

2015-01-15

**Valid until:**

2020-01-14

**Date,** 2015-01-16

Hans-Heiner Junker

TÜV SÜD Product Service GmbH is Notified Body with Identification no. 0423

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