



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 17 09 65725 019

Manufacturer:**Beijing Aeonmed Co., Ltd.**

11B2, Fengtai Science Park
100070 Beijing
PEOPLE'S REPUBLIC OF CHINA

EC-Representative:**Shanghai International Holding Corp. GmbH (Europe)**

Eiffestraße 80
20537 Hamburg
GERMANY

**Product Category(ies):**

**Anaesthetic Workstation, Vaporizer,
Ventilator, Medical Air Compressor,
Infusion Pump, Ceiling Pendant,
Medical Gas Terminal units,
Multi-Parameter Patient Monitor,
Videoscope System,
Patient Warming System.**

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.:

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Valid from:

2017-11-30

Valid until:

2020-05-03

Date, 2017-11-30

Stefan Preiß



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

Page 1 of 2



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No. G1 17 09 65725 019**Facility(ies):**

Beijing Aeonmed Co., Ltd.
11B2, Fengtai Science Park, 100070 Beijing, PEOPLE'S
REPUBLIC OF CHINA

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Hebei Province, PEOPLE'S REPUBLIC OF CHINA