



Product Service

EC Certificate

Production Quality Assurance

Directive 93/42/EEC on Medical Devices (MDD), Annex V
(Devices in Class IIa, IIb or III)

No. G2 11 10 78476 002

Manufacturer:**Xiamen Winner Medical Co., Ltd.**

4F, No.98 Huli Industrial Park
Mei Xi Dao
TongAn
361100 Xiamen
PEOPLE'S REPUBLIC OF CHINA

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

Eiffestraße 80
20537 Hamburg
GERMANY

Product**Category(ies):**

**Silicone/SEBS/PVC Manual Resuscitators
(Masks, Positive End Expiratory Pressure Valve,
Oxygen Tube, Reservoir), Resuscitation Mask,
Continuous Positive Airway Pressure
Mask/Non-Invasive Ventilation mask**

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

Report No.:

SH1171701

Valid from:

2019-12-13

Valid until:

2024-12-12



Hans-Heiner Junker

Date, 2019-12-15

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

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361100 Xiamen, PEOPLE'S REPUBLIC OF CHINA