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Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
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ZLG-BS-244.10.08



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 097063 0012 Rev. 00

Manufacturer: **Smiths Medical ASD, INC.**
6000 Nathan Lane North
Minneapolis MN 55442
USA

Product Category(ies): **Vibratory positive expiratory devices,
Incentive Spirometers, Silicone Laryngeal
Mask, Oxygen Line accessory,
Nasal Cannula, Oxygen Masks,
Drainage bags, Oxygen tubing,
Adult Face Tent, Disposable Anesthesia
Face Masks**

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. 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:G10970630012Rev.00

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Valid from: 2021-03-18
Valid until: 2024-05-26

Date, 2021-03-18

Christoph Dicks
Head of Certification/Notified Body