



Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zlg.de
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 010815 0038 Rev. 01

Manufacturer:

**Fisher & Paykel Healthcare Ltd.
15 Maurice Paykel Place
East Tamaki, Auckland 2013
NEW ZEALAND**

Product Category(ies): Respiratory Gas Delivery Systems,
Heated Humidifiers, Continuous Positive Airway
Pressure Units, Gas Powered Pulmonary Resuscitators,
Nasal and/or Oral Interfaces for Delivery of
Respiratory Gases, Patient Monitoring Software for
Use with Fisher & Paykel Healthcare Medical Devices,
Insufflation Gas Conditioning Systems

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.:

JAQ235040717

Valid from:

2019-12-12

Valid until:

2024-05-26

Date,

2019-12-12

Christoph Dicks
Head of Certification/Notified Body



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Facility(ies):

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15 Maurice Paykel Place, East Tamaki,
Auckland 2013, NEW ZEALAND**

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