



Product Service

EC - CERTIFICATE

Full Quality Assurance System

(Annex II, section 3 of the Directive 93/42/EEC on Medical Devices)

No. G1 10 09 20809 031

Manufacturer: **BREAS Medical AB**
 Företagsvägen 1
 43533 Mölnlycke
 SWEDEN

Facility(ies): BREAS Medical AB
 Företagsvägen 1, 43533 Mölnlycke, SWEDEN

Product Category(ies): **Respiratory Therapy Systems,
 Respiratory Monitoring Devices,
 Sleep Apnea and Humidifier 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 products / product categories according to Annex II section 3 of the Directive 93/42/EEC on Medical Devices. This quality assurance system conforms to the provisions of this Directive and is subject to periodical surveillance. For marketing of class III products an additional Annex II.4 certificate is mandatory. See also notes overleaf.

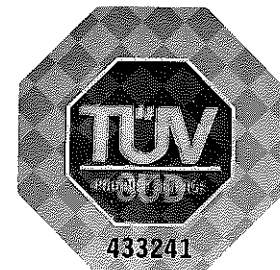
Report No.: 71373126

Valid until: 2015-10-27

H.-H.

Hans-Heiner Junker

Date, 2010-10-28



TÜV SÜD Product Service GmbH is Notified Body according to Council Directive 93/42/EEC concerning medical devices with identification no. 0123.

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