



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 15 11 93125 003

Manufacturer: **Amico Corporation**
85 Fulton Way
Richmond Hill ON L4B 2N4
CANADA



EC-Representative: **RMS-UK Limited**
28 Trinity Road
Nailsea
Somerset BS484NU
UNITED KINGDOM

Product Category(ies): **Medical Gas Outlets, Bed Head 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.

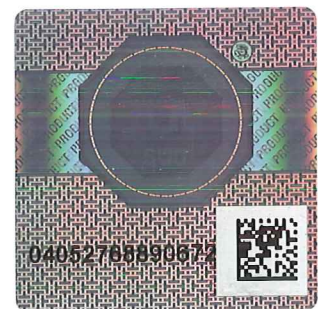
Report No.: 72110670

Valid from: 2015-12-28

Valid until: 2020-03-21

Date, 2015-12-29

Hans-Heiner Junker



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

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Facility(ies):

Amico Corporation
85 Fulton Way, Richmond Hill ON L4B 2N4, CANADA

Amico Corporation
120 Fulton Way, Richmond Hill ON L4B 1J5, CANADA