



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 10 05 62879 004

Manufacturer: Mediplus Pty Ltd
16 Redland Drive
Mitcham VIC 3132
AUSTRALIA

EC-Representative: MediMark Europe Sarl
11, Rue Emile Zola - BP 2332
38033 Grenoble Cedex 2
FRANCE

Product Category(ies): Breathing Circuits, Fittings and Adaptors

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.: ONQ73526982A

Valid until: 2012-06-11

Date, 2011-01-12

Hans-Heiner Junker



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

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Facility(ies):

**Mediplas Pty Ltd
16 Redland Drive, Mitcham VIC 3132, AUSTRALIA**