



Product Service

EC Certificate

Production Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex V
(Devices in class I in sterile conditions, sterilised systems or procedure packs)

No. G2S 16 03 11634 169

Manufacturer:

Medela AG

Lättichstrasse 4b
6341 Baar
SWITZERLAND



Facility(ies):

Medela AG
Lättichstrasse 4b, 6341 Baar, SWITZERLAND

Product Category(ies):

**Sterile suction tubing systems
and sterile canisters**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture in accordance with MDD Annex V. This quality assurance system covers those aspects of manufacture concerned with securing and maintaining sterile conditions of the respective devices / device categories and conforms to the requirements of this Directive. It is subject to periodical surveillance. See also notes overleaf.

Report No.:

713074241

Valid from:

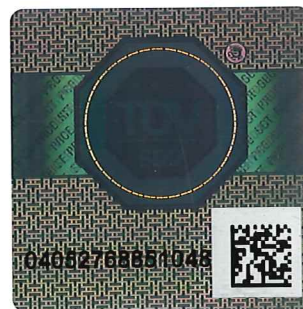
2016-04-18

Valid until:

2021-04-17

Date, 2016-04-12

Stefan Preiß



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

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