



Product Service

EC Certificate

Production Quality Assurance System

Directive 93/42/EEC on Medical devices (MDD), Annex V
(Devices in Class I with measuring function and in sterile condition)

No. G2MS 18 04 12974 456

Manufacturer:

B. Braun Melsungen AG

Carl-Braun-Str. 1
34212 Melsungen
GERMANY



Product Category(ies):

Sterile non-active medical devices for:
-Urology, gastroenterology, suction and drainage
as well as related configured customized sets
-Infusion systems for volume control incl.
accessories
-Measurement of central venous pressure
-Syringes, mixing containers
and dispenser sets for oral application

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for the manufacture in accordance with MDD Annex V. This quality assurance system covers those aspects of manufacture concerned with the metrological requirements and 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.:

713128919

Valid from:

2018-05-02

Valid until:

2023-05-01



Date, 2018-04-30

Stefan Preiß

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

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Facility(ies):

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