ONS AND





## **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 016329 0033 Rev. 00

Manufacturer

sfm medical devices GmbH

Brückenstraße 5 63607 Wächtersbach

**GERMANY** 

**Product** 

Sterile medical disposable cannulae,

Category(ies): syringes and filters

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.:

713157614

Valid from:

2020-03-13

Valid until:

2024-05-26

Date,

2020-03-13

Christoph Dicks

Head of Certification/Notified Body

Page 1 of 1 TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123