



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 18 04 84462 013

Manufacturer: **KARL STORZ SE & Co. KG**

Dr.-Karl-Storz-Straße 34
78532 Tuttlingen
GERMANY



Facility(ies): KARL STORZ SE & Co. KG
Dr.-Karl-Storz-Straße 34, 78532 Tuttlingen, GERMANY

Product Category(ies): **Sterile accessories of class I for endoscopic procedures**

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.: 713129927

Valid from: 2018-07-17

Valid until: 2023-07-16

Date, 2018-07-03

Stefan Preiß



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

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