



Product Service

EC Certificate

Full Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 16 04 27560 019

Manufacturer: **Fritz Ruck**
Ophthalmologische Systeme GmbH
 Ernst-Abbe-Str. 30b
 52249 Eschweiler-Dürwiß
 GERMANY



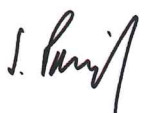
Facility(ies): Fritz Ruck Ophthalmologische Systeme GmbH
 Ernst-Abbe-Str. 30b, 52249 Eschweiler-Dürwiß, GERMANY

Product Category(ies): **Ophthalmological Surgical Systems**
including associated disposables

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.: 713078728

Valid from: 2016-04-05
Valid until: 2021-04-04

Date, 2016-04-01

 Stefan Preiß



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