



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 074222 0013 Rev. 01

Manufacturer: **Creagh Medical Ltd**
IDA Business Park
Ballinasloe, Co Galway, H53 K8P4
IRELAND

**Product Category(ies): PTA Catheters for Peripheral Applications
and Intravascular Microflow Catheters**

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. All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:G1074222_0013_Rev_01

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Valid from: 2021-05-18

Valid until: 2024-05-26

Date, 2021-05-18

Christoph Dicks
Head of Certification/Notified Body