EC Design-Examination Certificate

TÜVRheinland®

Directive 93/42/EEC on Medical Devices, Annex II (4)

Registration No.: ID 1191937-1

Manufacturer: Terumo Medical Corporation
950 Elkton Boulevard
Elkton MD 21921
USA

Products: Products Included:

Destination® Guiding Sheath
- carotid use (6 Fr, 7 Fr)
- renal use (5 Fr, 6 Fr, 7 Fr)
- peripheral use (5 Fr, 6 Fr, 7 Fr, 8 Fr)

The Notified Body hereby declares that an examination of the design dossier relating to the listed products has been performed according to Annex II, section 4 of the directive 93/42/EEC and that the design of the devices conforms to the requirements of the above-mentioned directive.

Report No.: 234171339-50

Effective date: 2021-05-06

Expiry date: 2024-05-26

Issue date: 2021-05-06

TÜV Rheinland LGA Products GmbH Zertifizierungsstelle TÜVRheinland®

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TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.

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