

EC Design-Examination Certificate



according to directive 93/42/EEC,
annex II (4)

As a notified body of the European Union, DEKRA Certification GmbH certifies for the manufacturer

JOTEC GmbH

Lotzenäcker 23, 72379 Hechingen, Germany

that the design dossier for the product(s) described in the annex complies with the requirements of the directive 93/42/EEC. This certificate is based on the result of the examination of the design dossier according to the directive 93/42/EEC annex II.4 as documented in the report mentioned in the Annex.

Product: E-tegra Stent Graft System

Date of the first certification: 2015-11-12

Date of the last recertification: ---.---.---

This certificate is valid until: 2020-11-11

Certificate registration No.: 50736-23-J0
Duplicate

DEKRA Certification GmbH
Stuttgart, 2015-11-12 Notified
Body ID-number: 0124



Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
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ZLG-BS-296.10.01

Lack of fulfilment on conditions as set out in the Certification Agreement may render this certificate invalid

