Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08

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 070145 0044 Rev. 02

Manufacturer:

William A. Cook Australia Pty. Ltd.
95 Brandl Street
Eight Mile Plains, Queensland 4113
AUSTRALIA

Product Category(ies):

Non-Active Medical Devices - Endovascular Grafts and Systems (Including Catheters); IVF Devices and Sets (Needles, Catheters, Tubing); Chorion Villus Biopsy Needle Sets;
Active Medical Devices - IVF Devices (Vacuum Pumps, Test Tube Heaters and Incubators)

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.: **JNQ235037629**

Valid from: **2020-04-24**
Valid until: **2024-05-26**

Date, **2020-04-24**

Christoph Dicks
Head of Certification/Notified Body

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

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

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