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

Production Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex V
(Devices in class I in sterile conditions, sterilised systems or procedure packs)

No. G2S 004384 0002 Rev. 02

Manufacturer

Curas Ltd
69-A, Raja Uda Business Centre Road 4
12300 Penang
MALAYSIA

Product Category(ies):

- **Sterile Prefilled Syringe**
- **Sterile Urine and Fluid Drainage Bags (Closed System Bags including TURB/P Bags)**
- **Sterile Luer Lock Caps**
- **Sterile Catheter Spigot**
- **Sterile Disposable Deep Trays and Bowls**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture in accordance with MDD Annex V. This quality assurance system covers those aspects of manufacture concerned with securing and maintaining sterile conditions of the respective devices / device categories and conforms to the requirements of this Directive. It is subject to periodical surveillance. See also notes overleaf.

Report No.: MYQMH0819043Rev1-721423522

Valid from: 2020-02-17
Valid until: 2024-01-27

Date, 2020-02-17

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

ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT
TÜV SÜD
A4 / 07.17

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Facility(ies):

Curas Ltd
69-A, Raja Uda Business Centre Road 4, 12300 Penang,
MALAYSIA

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