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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 101751 0002 Rev. 00

Manufacturer	**Centra For Medical Supplies Limited Partnership** Tower S 10 New FuStat City Cairo EGYPT
EC-Representative:	Obelis s.a 53, Boulevard General Wahis, 1030 Brussels, BELGIUM
Product Category(ies):	**Sterile Gowns and Sterile Drapes and Sterile Packs**

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.: IND2018042

Valid from: 2019-01-03
Valid until: 2024-01-02

Date, 2019-01-03

Stefan Preiß

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

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Facility(ies):

Centra For Medical Supplies Limited Partnership
Tower S 10 New FuStat City, Cairo, EGYPT

Centra For Medical Supplies Ltd
Garbia, Tanta - Kafr Elshikh Road, Qutor, EGYPT

-/-

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