EC Certificate Full Quality Assurance System: Certificate BE19/819943775

The management system of

Tobrix B.V.

Van Dijklaan 27
5581 WG Waalre, The Netherlands

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex II (excluding Section 4)

For the following products

Sterile catheters for endovenous laser ablation (peripheral applications)
Sterile baskets for kidney stone removal.
Sterile surgical laser fibers for use in laser surgery applications

Where the above scope includes class III medical device(s), a valid EC Design Examination Certificate according to Annex II (Section 4) is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 16 December 2019 until 11 May 2021
and remains valid subject to satisfactory surveillance audits.
Issue 1. Certified since 09 July 2013
and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered BE/AND 12/1312.QMD

Authorised by

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