

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 V

For the following products

**Sterile surgical Laser Probes for endovenous laser ablation,
Lithotripsy, Prostate ablation.**

For placing on the market of Class IIb or Class III devices covered by this certificate, an EC Type Examination Certificate according to Annex III is required.

This certificate is valid from 9 January 2015 until 11 May 2018 and
remains valid subject to satisfactory surveillance audits.

Re certification audit due before 11 April 2016.

Issue 2. Certified since 9 July 2013.

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

Authorised by

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