EC Certificate Full Quality Assurance System: Certificate US00/52267

SGS

The management system of

Microline Surgical, Inc.

50 Dunham Road, Suite 1500

Beverly, MA, 01915. United States 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

Active laparoscopic surgical instruments, sterile and non-sterile scissor, grasping and dissecting tips; sterile and non-sterile clip appliers, sterile ligating clips; sterile cautery forceps, sterile and non-sterile thermal ligation shears (including power supply and footswitch); port access system.

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 14 August 2015 until 30 July 2020 and remains valid subject to satisfactory surveillance audits.

Re certification audit due before 29 July 2018 Issue 14. Certified since 12 December 2000 Certification is based on reports numbered WW/ME 201551

Authorised by



SGS United Kingdom Ltd, Notified Body 0120

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