EC Certificate Full Quality Assurance System: Certificate KR11/01589

The management system of

Sung Hwan E&B Co.,Ltd.

#502, SK Techno Bldg, 16-4 Seongsu-dong 1-ga, Seongdong-gu,
Seoul, Korea

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

**High Frequency stimulator (Model : SH-312B, SH-304);
High frequency electrosurgical unit (model: VIVACE) including associated hand piece, foot switch and sterile micro needle cartridge(model: SH-TIP)**

For placing on the market of Class III devices covered by this certificate, an EC Design Examination Certificate according to Annex II (Section 4) is required.

This certificate is valid from 22 May 2014 until 22 August 2014 and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 21 January 2014
Issue 5. Certified since 22 February 2011

Certification is based on reports numbered KR/SEL Y-PC/10250

Authorised by

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