EC Certificate Full Quality Assurance System: Certificate US14/841851

The management system of

Kirwan Surgical Products LLC

180 Enterprise Drive, Marshfield, MA, 02050, 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

Sterile and Non-sterile electrosurgical handpieces, bipolar coagulators and Handpieces with Cords.

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 08 July 2019 until 24 May 2024
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 07 March 2022
Issue 4. Certified since 09 June 2013

Certification is based on reports numbered WW/MW 606189

Authorised by

Jonathan M. Hall

SGS United Kingdom Ltd, Notified Body 0120
202B Worle Parkway, Weston-super-Mare, BS22 6WA UK
t +44 (0)1934 522917 f +44 (0)1934 522137 www.sgs.com

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