

EC Certificate Full Quality Assurance System: Certificate GB08/75138

The management system of

Finesse Medical Limited

Unit 4, Royal Canal Business Park,
Athlone Road, Longford, Ireland

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

Polyurethane foam wound dressings, Silver alginate catheter dressings, Sterile hydrogel wound dressings, Sterile skin barrier film, Sterile Elta Silfoam and Silfoam Border wound dressings. Tracheostomy wound dressing.

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 07 July 2011 until 07 July 2016 and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 19 June 2014
Issue 7. Certified since 07 July 2008

Certification is based on reports numbered GB/PC 216465

Authorised by

SGS United Kingdom Ltd, Notified Body 0120

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