EC Certificate Production Quality Assurance System: Certificate US06/70137

The management system of

AngioSystems, Inc.

7 Hopkins Place,
Ducktown, TN, 37326, United States

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 procedure-ready trays and kits (including disposable syringes, needles, instruments, and gauze).

Where the above scope includes class IIb or class III medical device(s), a valid EC Type Examination Certificate according to Annex III is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 4 February 2016 until 17 October 2020
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 12 September 2018
Issue 4. Certified since 17 October 2006

Certification is based on reports numbered WW/ME 214284

Authorised by

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