EC Certificate Production Quality Assurance System: Certificate GB00/51465

The management system of

Wescott Medical Ltd

Unit 3B, Drum Industrial Estate, Chester-le-Street, Durham, DH2 1AG, UK

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex V

For the following products

Sterile infusion sets and infusion extension sets; Infusion connectors valves filters and caps; syringe – pump extension sets; medical gas tubing sets.

Sterility aspects only - Restricted to the aspects of manufacture concerned with securing and maintaining sterile conditions: Irrigation sets and systems.

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 25 August 2015 until 20 March 2020
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 20 March 2018
Issue 10. Certified since 20 March 2000

Certification is based on reports numbered GB/PC 08905

Authorised by

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