

EC Certificate Production Quality Assurance System: Certificate
GB98/14492

The management system of

Casmed International Ltd

Unit C, Blenheim House, 1 Blenheim Road,
Epsom, Surrey, KT19 9AP, UK

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex V

For the following products

The scope of registration appears on page 2 of this certificate

This certificate is valid from 07 September 2013 until 07 September 2018
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 07 September 2016
Issue 11. Certified since 07 September 1998

Certification is based on reports numbered GB/PC 09152

Authorised by



SGS United Kingdom Ltd, Notified Body 0120

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SGS CE 13 0311 M2

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Casmed International Ltd

Directive 93/42/EEC

on medical devices, Annex V

Issue 11

Detailed scope

Directive 93/42/EEC
on medical devices, Annex V

Sterile and non-sterile trocar and cannula, tubing sets, oocyte collection devices, procedure needles and needle guides, embryo transfer catheters, intra uterine insemination catheters.

Annex V Sterility aspects only - Restricted to the aspects of manufacture concerned with securing and maintaining sterile conditions

Sterile camera drapes.

For placing on the market of Class IIb or Class III devices covered by this certificate, an EC Type Examination Certificate according to Annex III is required.