EC Certificate Full Quality Assurance System: GB98/13044

The management system of

Research Instruments Ltd

Bickland Industrial Park, Falmouth, Cornwall, TR11 4TA, UK

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

The scope of registration appears on page 2 of this certificate.

This certificate is valid from 30 October 2018 until 05 August 2021
andremains valid subject to satisfactory surveillance audits.
Re certification audit due before 07 July 2019
Issue 18. Certified since 29 May 1998

Certification is based on reports numbered GB/PC 08973

This is a multi-site certification.
Additional site details are listed on the subsequent page.

Authorised by

SGS United Kingdom Ltd, Notified Body 0120

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SGS CE 02 0315 M2

Research Instruments Ltd

Directive 93/42/EEC

on medical devices, Annex II (excluding section 4)

Issue 18

Detailed scope

Integra micromanipulation system, Saturn laser systems, electrically heated plates and temperature control units for use in RI Witness systems, sterile single-use glass RI pipettes, sterile single-use plastic pipettes (EZ Range) and sterile single-use migration sedimentation chambers (RI MSC) all for use in handling and treatment of human reproductive samples.

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

Additional facilities

Site B, Bickland Industrial Park, Falmouth, Cornwall, TR11 4TA, UK

Site C, Bickland Industrial Park, Falmouth, Cornwall, TR11 4TA, UK