EC Certificate Full Quality Assurance System: Certificate GB19/964592

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 16 December 2019 until 24 May 2024
and remains valid subject to satisfactory surveillance audits.
Issue 1. Certified since 29 May 1998
and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered GB/PC 08973

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

Authorised by

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Research Instruments Ltd

Directive 93/42/EEC

on medical devices, Annex II (excluding Section 4).

Issue 1

Detailed scope

Integra micromanipulation system, Saturn laser systems, electrically heated plates and temperature control units f or 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