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

SGS Belgium NV, Notified Body 1639

SGS House Noorderlaan 87 2030 Antwerp Belgium t +32 (0)3 545-48-48 f +32 (0)3 545-48-49 www.sgs.com

LPMD5007 - Certificate CE1639 Annex II-4_EN rev. 02

Page 1 of 2



This document is issued by the Company subject to its General Conditions of Certification Services, unless otherwise agreed, accessible at www.sgs.com/terms_and_conditions.htm. Altention is drawn to the limitations of liability, indemnification and jurisdictional issues established therein. The authenticity of this document may be verified at https://www.sgs.com/en/certified-clients-and-products/certified-client-directory. Any unauthorized alteration, forgery or falsification of the content or appearance of this document is unlawful and offenders may be prosecuted to the fullest extent of the law.



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.

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