

Medical Device Full Quality Assurance System Certificate GB23/00000053

The management system of

Planer Limited

110 Windmill Road Sunbury-on-Thames Middlesex TW16 7HD United Kingdom

has been assessed and certified as meeting the requirements of
**Part II of The Medical Devices Regulations 2002, Annex II excluding
section 4 [as modified by Part 2 of Schedule 2A to The Medical
Devices Regulations 2002]**

For the following products

Incubators intended for assisted reproductive technologies:

Incubators:

BT37

Where the above scope includes class III medical device(s), a valid Design Examination Certificate according to Annex II (Section 4) [as modified by Part 2 of Schedule 2A of The MDR 2002] is a mandatory requirement for each device in addition to this certificate to place that device on the market

Certification is based on reports numbered GB/PC/240639

Previous certificate number: N/A

Change in between this certificate and previous one: No changes from previous certification scope

This certificate is valid from 12 May 2026 until 12 May 2031 and remains valid subject to satisfactory surveillance audits.

Issue 4. Certified since 31 January 2023



Authorised by

Robert Snell

Head of Approved Body

SGS United Kingdom Ltd Approved Body 0120

Rossmore Business Park, Ellesmere Port, Cheshire, CH65 3EN, UK

t +44 (0)151 350-6666 - www.sgs.com

This document is an authentic electronic certificate for Client' business purposes use only. Printed version of the electronic certificate are permitted and will be considered as a copy. This document is issued by the Company subject to SGS General Conditions of certification services available on [Terms and Conditions | SGS](#). Attention is drawn to the limitation of liability, indemnification and jurisdictional clauses contained therein. This document is copyright protected and any unauthorized alteration, forgery or falsification of the content or appearance of this document is unlawful.

