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

R Snell

Authorised by
Robert Snell
Head of Approved Body

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