Medical Device Full Quality Assurance System Certificate GB23/00000235

The management system of

Lynton Lasers Ltd

Lynton House Manor Lane Holmes Chapel Cheshire CW4 8AF 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
LUMINA and 3JUVE/EXCELIGHT Intense Pulsed Light and Laser Systems for the treatment of superficial vascular lesions, scars, superficial pigmentation removal, active acne control, and the removal of unwanted hair when being carried out as part of the treatment for Polycystic Ovary Syndrome, pilonidal sinus or pseudofolliculitis barbae.

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/07602

Previous certificate number: N/A

Change in between this certificate and previous one: N/A

This certificate is valid from 01 June 2023 until 04 March 2024 and remains valid subject to satisfactory surveillance audits.
Issue 1. Certified since 01 June 2023

Authorised by
Lynsey Hall
Head of Approved Body 0120

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