Medical Device Full Quality Assurance System Certificate GB23/0000007

The management system of

Alesi Surgical Limited

Cardiff Medicentre Heath Park Cardiff CF14 4UJ 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

Ultravision Electrostatic Precipitator and sterile trocars for use in surgical procedures.

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/230872

Previous certificate number: N/A

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

This certificate is valid from 05 January 2023 until 05 January 2028 and remains valid subject to satisfactory surveillance audits.

Issue 1. Certified since 05 January 2023

Authorised by

Lynsey Hall

Head of Approved Body 0120

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