The management system of

LaproSurge Limited

5b Fishers Industrial Estate, Wiggenhall Road, Watford, Hertfordshire, WD18 0FN, 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

Sterile Laparoscopic Instruments for single use:

Sterile Monopolar Laparoscopic Scissors, Graspers & Dissectors

Sterile Disposable Diathermy Electrodes

Sterile Bipolar disposable Maryland Forceps

Sterile suction Irrigation sets with irrigation/aspiration probes

with or without Diathermy

Sterile Insufflation Needle

Sterile Tissue retrieval system

Sterile Insufflation tubing

Sterile Endoscopic Anti-Fog Solution with pad

Sterile Trocar & Cannula and Trocar Site Closure Device

Sterile Disposable laparoscopic Blunt Tipped Dissectors

Where the above scope includes class III medical device(s), a valid EC Design Examination 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.

This certificate is valid from 01 April 2021 until 24 May 2024 and remains valid subject to satisfactory surveillance audits.

Issue 5. Certified since 01 October 1999.

Certification is based on reports numbered GB/PC 06917

Authorised by

Global Medical Devices Head of Notified Body

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 1



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