Medical Device Production Quality Assurance System Certificate
GB23/00000253

The management system of

Farla Medical Healthcare Limited

Unit 2 1000 North Circular Road London NW2 7JP United Kingdom

has been assessed and certified as meeting the requirements of

Part II of The Medical Devices Regulations 2002, Annex V [as modified by Part 2 of Schedule 2A to The Medical Devices Regulations 2002]

For the following products

Sterile single use surgical procedure packs and surgical instruments
(for gynaecology, ophthalmology podiatry ENT, and general surgery).
Sterile surgical sampling blades.
Sterile Biopsy punch.
Sterile lubricating jelly.

Annex V - sterility aspects only - restricted to the aspects of manufacture concerned with securing and maintaining sterile conditions:

Sterile single use non surgically invasive & non active instruments and procedure packs:
Sterile Gynaecological collector (brush).
Sterile single use instruments for gynaecology, ophthalmology, podiatry,
ENT and general surgery
Sterile dressing packs.
Sterile dressings and drapes.

Where the above scope includes class IIb or class III medical device(s), a valid Type Examination Certificate according to Annex III [as modified by Part 2 of Schedule 2A to 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/216926

Previous certificate number: N/A

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

This certificate is valid from 22 June 2023 until 24 May 2024 and remains valid subject to satisfactory surveillance audits.
Issue 1. Certified since 22 June 2023

L. Henderson

Authorised by
Lynn Henderson

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