EC Certificate Full Quality Assurance System: Certificate PK19/818842594

The management system of

SURGIKARE

Toor Abad Daska Road, Sialkot, Pakistan

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 & Non-Sterile Single Use Bipolar Forceps with & Without Cables, Sterile & Non-Sterile Single Use Bipolar Electrodes, Non-Sterile Reusable Bipolar Forceps with & without cables, Non-Sterile Reusable Bipolar Electrodes.

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 16 December 2019 until 29 June 2023
and remains valid subject to satisfactory surveillance audits.
Issue 1. Certified since 25 November 2019
and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered PK/LHR 221615

Authorised by

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