EC Certificate Production Full Quality Assurance System: Certificate PK19/818842593

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 V

For the following products

Sterile and non-sterile single-use surgical instruments: forceps, scissors, retractors, tweezers, curettes, cannulae, suction tubes, needle holders, bone cutters, probes, trocars, scalpel handle without blade, Sterile single-use catheters used for drainage of body fluids

Annex V Sterility Aspects Only - restricted to the aspects of manufacture concerned with securing and maintaining sterile conditions:

Sterile single-use devices: dental scalars, extracting forceps, mouth gags, protoscopes, dental syringes without needle, dilators, vaginal speculae, dental elevators.

Where the above scope includes class IIb or class III medical device(s), a valid EC Type Examination Certificate according to Annex III is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 15 April 2021 until 29 June 2023
and remains valid subject to satisfactory surveillance audits.
Issue 2. Certified since 29 June 2012

Certification is based on reports numbered PK/LHR/ 221615

Authorised by

Global Medical Devices Head of Notified Body

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