

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

The scope of registration appears on page 2 of this certificate.

This certificate is valid from 16 December 2019 until 29 June 2023
and remains valid subject to satisfactory surveillance audits.

Issue 1. Certified since 29 June 2012
and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered PK/LHR/ 221615

Authorised by

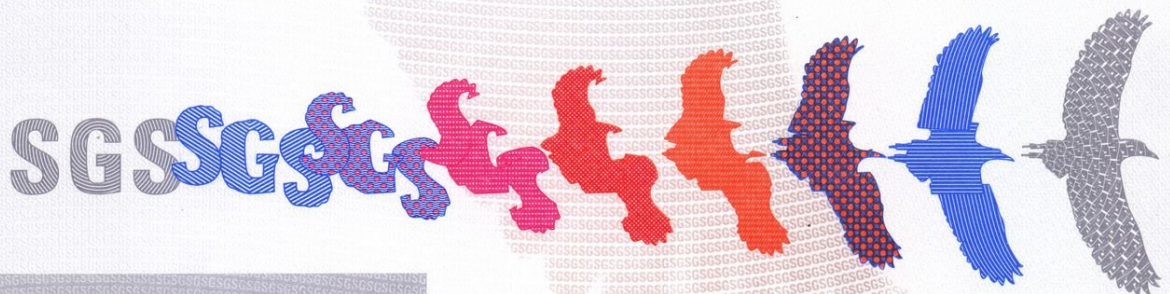
Pieter Weterings
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LPMDS008 - Certificate CE1639 AnnexV_EN rev. 01

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FH 0511

SURGIKARE

Directive 93/42/EEC

on medical devices, Annex V

Issue 1

Detailed scope

**Sterile & Non sterile single use surgical Forceps,
Sterile & Non sterile single use surgical scissors,
Sterile & Non Sterile single use surgical Retractors,
Sterile & Non Sterile single use Surgical Tweezers,
Sterile & Non sterile single use surgical curettes,
Sterile & Non sterile single use Surgical Cannulas,
Sterile & Non sterile single use Suction Tubes,
Sterile & Non sterile Single use Needle Holders,
Sterile & Non sterile single use Bone Cutters,
Sterile & Non Sterile Single Use Surgical Probes,
Sterile & Non sterile single use Trocars,
Sterile & Non sterile single use 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 Dental Scalars,
Sterile Single use Extracting Forceps,
Sterile Single use Mouth Gag,
Sterile single use Proto scopes,
Sterile Single Use Dental Syringes without needle,
Sterile Single use Dilators,
Sterile single use Vaginal Speculums,
Sterile single use Dental Elevator**

