EC Certificate Full Quality Assurance System: Certificate PK19/818842624.02

The management system of

Cosil Instruments Company

0.25 Km Daska Road, Ghuinke, 51040 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 Cables,
Sterile & Non Sterile Single Use Electrosurgical Pencils with Electrodes,
Non Sterile Reusable Electrosurgical Pencils with Electrodes,
Non Sterile Reusable Bipolar Forceps
with Cables and Non Sterile Reusable Monopolar Forceps with Cables.**

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 24 May 2024
and remains valid subject to satisfactory surveillance audits.
Issue 1. Certified since 05 June 2013
and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered PK/LHR 230240

Multiple certificates have been issued for this scope.
The main certificate is numbered PK19/818842624.00

Authorised by

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

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