

The management system of

Tecno Instruments Pvt. Ltd.

316-C Small Industrial Estate Sialkot - 51340, 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

Bipolar Forceps Reusable, Monopolar Forceps / Diathermy Inst Reusable, Bipolar Forceps sterile Single use w/wo cable, Monopolar Forceps Sterile Single use w/wo cable, Finger Switch Pencil Reusable, Finger Switch Pencil sterile single use, Electrosurgical Electrodes Reusable, Electrosurgical Electrodes Sterile Single use, Non Stick Bipolar Forceps Reusable, Bipolar Artery Sealer Reusable, Bipolar Scissor Reusable

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 13 March 2023 and remains valid subject to satisfactory surveillance audits.

Issue 1. Certified since 20 August 2000 and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered PK/LHR 201127

Authorised by

SGS Belgium NV, Notified Body 1639

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LPMD5007 - Certificate CE1639 Annex II-4_EN rev. 02

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