

EC Certificate Full Quality Assurance System: Certificate PK19/818842627

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

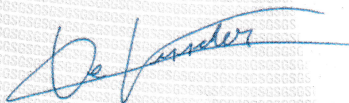
Where the above scope includes class III medical device(s), a valid EC Design Examination Certificate according to Annex II (Section 4) is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 25 March 2021 until 24 May 2024 and remains valid subject to satisfactory surveillance audits.

Issue 2. Certified since 20 August 2000

Certification is based on reports numbered PK/LHR 201127

Authorised by



Global Medical Devices Head of Notified Body

SGS Belgium NV, Notified Body 1639

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