

EC Certificate Full Quality Assurance System: Certificate GB00/51862

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 27 July 2015 until 13 March 2020 and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 22 January 2018
Issue 9. Certified since 29 August 2000

Certification is based on reports numbered GB/PI 201127

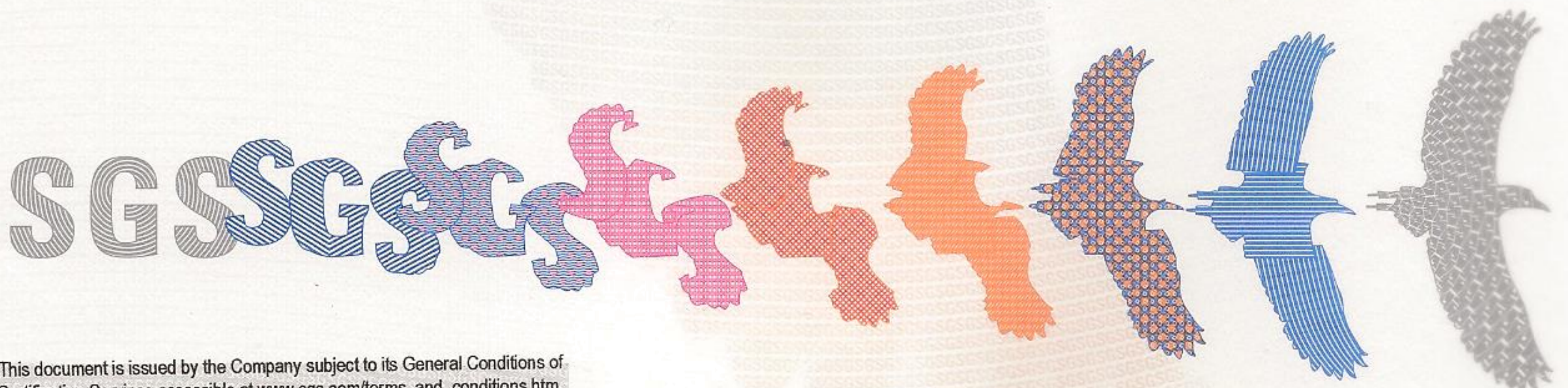
Authorised by

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