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

Reusable Non Sterile Bipolar Forceps complete with Electrosurgical Cables, Reusable Non Sterile Monopolar Forceps complete with Electrosurgical Cables and Reusable Non Sterile Finger Switch Pencils complete with Electrodes.

Sterile Single Use Bipolar Forceps complete with Electrosurgical Cables, Sterile Single Use Monopolar Forceps complete with Electrosurgical Cables and Sterile Single Use Finger Switch Pencils complete with Electrodes.

For placing on the market of Class III devices covered by this certificate, an EC Design Examination Certificate according to Annex II (Section 4) is required.

This certificate is valid from 13 March 2012 until 13 March 2017 and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 22 January 2015
Issue 6. Certified since 29 August 2000

Certification is based on reports numbered GB/PI 201127

Authorised by

SGS United Kingdom Ltd, Notified Body 0120

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