EC Certificate Production Quality Assurance System: Certificate PK15/92789

The management system of

Beauty Teck International (Pvt) Ltd.

Head Marala Road, Sialkot 51310 - Pakistan.
has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex V

For the following products

Non-sterile single use Scissors, Non-sterile single use Artery Forceps, Non-sterile. single use Dressing & Tissue Forceps, Non-sterile single use Surgical Curettes, Non-sterile single use Needle Holders, Non-sterile single use Bone Roungers, Non-sterile single use Cannulas for Injecting Medicine in Vein, Non-sterile single use Scalpel Handle & Knife and Non-sterile single use Retractors

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

This certificate is valid from 06 March 2018 until 06 March2023
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 02 December 2021
Issue 2. Certified since 06 March 2015

Certification is based on reports numbered GB/PI 233480

Authorised by

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