

The management system of

Intelligent Medical Innovations

6 km, Daska Road,
Sialkot, 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

GERATI single use sterile electrosurgical dissecting forceps, GERATI single use sterile electrosurgical scissors, GERATI single use sterile non powered grasping forceps, GERATI single use sterile non powered tissue holding forceps.

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 09 August 2011 until 09 August 2014 and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 25 July 2014
Issue 1. Certified since 09 August 2011

Certification is based on reports numbered GBPI 225468

Authorised by

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Page 1 of 1

