

The management system of

Afri Medical Company

13, Obour Buildings, Salah Salem Street,
Box 215, Heliopolis, Cairo, Egypt

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex II (excluding Section 4) and Annex V

For the following products

The scope of registration appears on page 2 of this certificate.

This certificate is valid from 26 February 2011 until 10 January 2014 and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 10 January 2014
Issue 7. Certified since 10 January 2005

Certification is based on reports numbered GB/P1 211410

This is a multi-site certification.
Additional site details are listed on the subsequent page.

Authorised by

SGS United Kingdom Ltd, Notified Body 0120

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Afri Medical Company

Directive 93/42/EEC

on medical devices, Annex II (excluding section 4) and Annex V

Issue 7

Detailed scope

Annex II (excluding Section 4)

Dialysers.

Annex V

**Blood line, Fistula needle, infusion sets, transfusion sets,
Endotracheal tubes, Suction catheters, sterile oxygen masks, sterile
irrigators, Sterile Surgical Dressing and sterile Incise Drape.**

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.

Additional facilities

Industrial Area C3, 10th of Ramadan City, Egypt