

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 V

For the following products

The scope of registration appears on page 2 of this certificate

For Placing on the market of class IIb or class III devices covered by this certificate, an EC
type Examination according to annex III is required

This certificate is valid from 8 July 2015 until 10 January 2019 and remains
valid subject to satisfactory surveillance audits.

Re certification audit due before 10 January 2017

Issue 3. Certified since 10 January 2005

This is a multi-site certification.

Additional site details are listed on the subsequent pages.

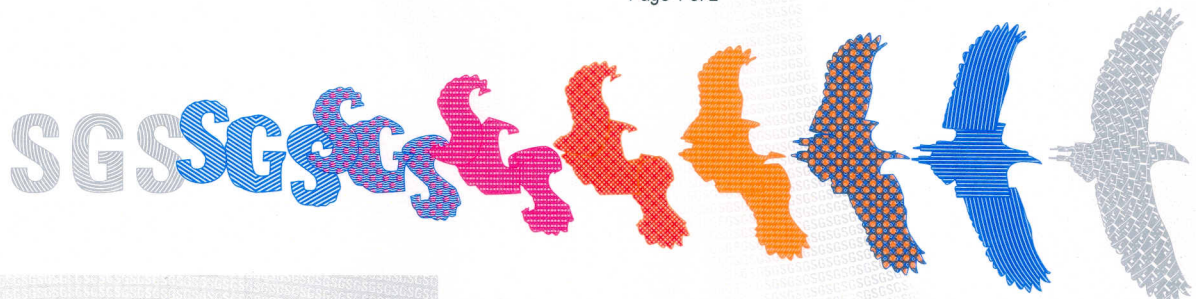
Authorised by

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Afri Medical Company

Directive 93/42/EEC

on medical devices, Annex V

Issue 3

Detailed scope

**Blood line, Fistula needle, Sterile Surgical Dressing and sterile
Incise Drape.**

For Placing on the market of class IIb or class III devices covered by this certificate, an EC
type Examination according to annex III is required

Additional facilities

**Industrial Area C3, 10th of Ramadan City, Egypt , 10th of Ramadan,
Egypt**