EC Certificate Production Quality Assurance System: Certificate
EG05/63842

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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SGS CE 14 0315 M2

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