EC Certificate Full Quality Assurance System: EG14/3203

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)

For the following products

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

This certificate is valid from 13 March 2014 until 10 January 2019 and
remains valid subject to satisfactory surveillance audits.
Re certification audit due before 10 January 2017
Issue 1. Certified since 10 January 2005

Certification is based on reports numbered PI 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
SGS United Kingdom Ltd Systems & Services Certification
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SGS CE 01 0311 M2

Afri Medical Company

Directive 93/42/EEC

on medical devices, Annex II (excluding section 4)

Issue 1

Detailed scope

Dialysers

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 , 10th of Ramadan, Egypt