EC Certificate Production Quality Assurance System: Certificate EG19/19020

The management system of

Farcomake For Advanced Medical Industry

New Borg Al-Arab Industrial City, Area 4, Block 2 Part 7, 16
Alexandria, Egypt

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex V

For the following products

Annex V Sterile IV Cannula, Sterile Bloodlines and Sterile Fistula needle, sterile endotracheal tube, sterile syringes with needle, sterile infusion set.
Annex V Sterility aspects only - Restricted to the aspects of manufacture concerned with securing and maintaining sterile conditions sterile 3-way stop cock syringe without needle sterile extension tube

Where the above scope includes class IIb or class III medical device(s), a valid EC Type Examination Certificate according to Annex III is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 16 December 2019 until 18 April 2023
and remains valid subject to satisfactory surveillance audits.
Issue 1. Certified since 21 November 2008
and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered EG/CAI/ PI219250

Authorised by

SGS Belgium NV, Notified Body 1639
SGS House Noorderlaan 87 2030 Antwerp Belgium
t +32 (0)3 545-48-48 f +32 (0)3 545-48-49 www.sgs.com