EC Certificate Production Quality Assurance System: Certificate EG20/2833

The management system of

Euromed for Medical Industries (S.A.E)

Area (10) Block (i), Free zone Nasr City, 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

Sterile Infusion Sets (Burette), Sterile Syringes, Sterile Tracheal Tubes, Sterile IV Cannula, Sterile Foley Catheters, Sterile Silicone Foley Catheters, Sterile infusion Set, Disposable Sterile Needle And Sterile Safety Syringes Annex V (Sterility Aspects Only) - Restricted To The Aspects Of Manufacture Concerned With Securing And Maintaining Sterile Conditions Sterile 3-Way Stopcock

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 28 October 2020 until 24 May 2024
and remains valid subject to satisfactory surveillance audits.
Issue 4. Certified since 25 October 2006
and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered EG/CAI/ PI215191

Authorised by

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