EC Certificate Production Quality Assurance System: Certificate PK19/818842587

The management system of

Qatari German for Medical Devices Company

Building No. 136, Street 54, Abu Hamour, P.O.Box 22556, Doha-Qatar

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex V

For the following products

Q JECT - Sterile Single Use Hypodermic Syringes with & without needle.
Q JECT Ultra - Sterile Single Use Syringe for Insulin Use
Q SAFE - Sterile Single Use Safety Syringes with Re-Use Prevention Features (RUP)
Q FLOW - Intravacular Catheter or Cannula
Q NEED - Sterile Single Use Hypodermic Needles

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 04 April 2024
And remains valid subject to satisfactory surveillance audits.
Issue 1. Certified since 10 April 2019
and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered PK/LHR/ PK18357

Authorised by

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LPMD5008 - Certificate CE1639 AnnexV_EN rev. 01