EC Certificate Full Quality Assurance System: Certificate TW19/20064

The management system of

Lily Medical Corporation

No.28-2, Shun Jeau Diann, Chunan Town
Miaoli County, 35056, Taiwan, R.O.C.

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

Restricted to the aspects of manufacture concerned with securing and maintaining sterile conditions, and the conformity of the devices with metrological requirements

Sterile single use Intravascular administration set including Infusion set, Volume Metric Administration set, Injection cap and Neutral Pressure Needleless Connector.

Where the above scope includes class III medical device(s), a valid EC Design Examination Certificate according to Annex II (Section 4) is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 09 October 2020 until 21 August 2023
and remains valid subject to satisfactory surveillance audits.
Issue 2. Certified since 21 August 2012
and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered TW/TPE K605096

Authorised by

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