EC Certificate Full Quality Assurance System: Certificate TW12/11825

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

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 15 November 2018 until 21 August 2023
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 11 July 2021
Issue 4. Certified since 21 August 2012

Certification is based on reports numbered TW/TPE K605096

Authorised by

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