EC Certificate Full Quality Assurance System: Certificate GB98/12498

The management system of

LiDCO Ltd

16 Orsman Road, London, N1 5QJ, UK

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

Hemodynamic Monitoring Equipment including software and sensor interface. CNAP (non-invasive blood pressure device for use with LiDCO Hemodynamic Monitoring Equipment). Blood Pressure Module (for use with LiDCO Hemodynamic Monitoring Equipment) Flow Regulator (for use with LiDCO Hemodynamic Monitoring Equipment). Sterile Injectate Kit.

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 20 May 2017 until 20 May 2022
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 28 April 2020
Issue 13. Certified since 13 March 1998

Certification is based on reports numbered GB/PC 08524

Authorised by

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