EC Certificate Full Quality Assurance System: CN14/10399

The management system of

Beijing KellyMed Co., Ltd.

Room 115, 4# Building, No.2 Tiyuan West Road, Haidian District,
Beijing, 100084, P.R. China

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

The scope of registration appears on page 2 of this certificate.

This certificate is valid from 3 July 2017 until 2 July 2022
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 1 August 2020
Issue 2. Certified since 23 September 2014

Certification is based on reports numbered CN/BJS 8004

This is a multi-site certification.
Additional site details are listed on the subsequent page.

Authorised by

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SGS CE 02 0315 M2

Beijing KellyMed Co., Ltd.

Directive 93/42/EEC

on medical devices, Annex II (excluding section 4)

Issue 2

Detailed scope

Infusion Pump used for intravenous infusion administration, Syringe Pump used for intravenous injection administration, Feeding Pump used to supply nutrition

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

Additional facilities

2nd Floor, No.1 Building, No.2 Jingshengnan Street #15, Jinqiao Industrial Base, Zhongguancun Science Park Tongzhou Sub-Park, Tongzhou District, 101102, Beijing, 101102, P.R. China