EC Certificate Full Quality Assurance System: Certificate CN19/41114

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 22 September 2020 until 24 May 2024
and remains valid subject to satisfactory surveillance audits.
Issue 2. Certified since 23 September 2014
and first certified by SGS Belgium NV since 16 December 2019.

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

Certification is based on reports numbered CN/BJS 8004

Authorised by

SGS Belgium NV, Notified Body 1639
SGS House Noorderlaan 87 2030 Antwerp Belgium
t +32 (0)3 545-48-48 f +32 (0)3 545-48-49 www.sgs.com

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, Beijing, 101102, P.R. China