EC Declaration of Conformity

Manufacturer:

Beijing KellyMed Co., Ltd.
Room 115,4# Building, No.2 Tiyuan West Road, Haidian District, Beijing, 100084, China

whose single Authorized Representative:

Labcon GmbH
Benzstrasze 4,64646 Heppenheim
Germany
Tel:0049 6252 9425 0
Email:Labcon@Labcon.de
DE0000046575

We, the manufacturer, herewith declare that the products

Syringe pumps
Model: KL-602，KL-702，KL-605T
*(*including system components and accessories*)*
GMDN Code: 13217

meet the provisions of Directive 93/42/EEC which apply to them.

The medical device has been assigned to class IIb according to Annex IX of the Directive 93/42/EEC. It bears the mark

CE 0120

The product concerned has been designed and manufactured under a quality management system according to Annex II of Directive 93/42/EEC.

Compliance of the designated product with the Directive 93/42/EEC has been assessed and certified by the Notified Body

SGS United Kingdom Ltd,
202B Worle Parkway, Weston-super-Mare,BS22 6WA UK

Certificate No.: CN14/10399
Issue date: 3 July 2017
Expiry date: 2 July 2022

following the procedure relating to the EC Declaration of Conformity set out in Annex II of Directive 93/42/EEC.

The above mentioned declaration of conformity is exclusively under the responsibility of

Beijing KellyMed Co., Ltd.

Beijing, 31 July 2017
Place , date

北京科力建元医疗科技有限公司 BEIJING KELLYMED CO., LTD.

Legally binding signature, Function