EC Declaration of Conformity

Manufacturer:

CDM Center of Excellence Ltd.
Golden Land Building Office 1302,
Liangmaqiao Road No.32, Chaoyang District, Beijing, 100125, China

UK Authorised Representative:

WellKang Ltd (www.CE-marking.eu)
Enterprise Hub, NW Business Complex,
1 Beraghmore Road,Derry, BT48 8SE,
Northern Ireland

We, the manufacturer, herewith declare that the products

Disposable Syringe Sets

*(*including system components and accessories*)*

UMDNS-Code: **13-929**

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

The medical device has been assigned to **Class IIa Rule 6 a**ccording to Annex IX of the Directive 93/42/EEC. It bears the mark

CE 0197

The product concerned has been manufactured under a quality management system according to Directive 93/42/EEC.

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

TÜV Rheinland LGA Product GmbH
Tillystraße2 - 90431 Nürnberg
Certificate No.: DD 60146829 0001
Issue date: February.11, 2020
Expiry date: May.26, 2024

following the procedure relating to the EC Declaration of Conformity set out in Annex VII coupled with Annex V of Directive 93/42/EEC.

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

CDM Center of Excellence Ltd.



Beijing 2021-10-11
Place and Date

Name/Signature of the company's Representative