EC Declaration of Conformity

Manufacturer:
Jiangsu Dynamic Medical Technology Co., Ltd
No.108 Xingpu Road, Lujia Town, Kunshan, City, Jiangsu Province, 215331.
PEOPLE' S REPUBLIC OF CHINA

whose single Authorized Representative:
Shanghai International Holding Corp Gmbh (Europe)
Eiffestrasse 80, 20537 Hamburg, Germany

We, the manufacturer, herewith declare that the products

Medical Suction Unit

Model: DS series.
DS1000、DS2501M、DS3701M、DS3701CS、DS3701CS-1、DS3701CS-2011、DS7501CS、DS01、DS02、DS03、DS04、DS06、DS08、DS10、DS301、DS302、DS502、DS504、DS506、DS508、DS510、DAS001、DAS002、DS803、DS805、DS808、DS810、DS910、DS915、DS920、DS930、DS940、DS950

UMDNS-Code: 10-212

meet the provisions of Directive 93/42/EEC which apply to them.
The medical device has been assigned to class I (active) according to Rule 12 of Annex IX of the Directive 93/42/EEC. It bears the mark

CE

The product concerned has been manufactured according to Annex V.3 of Directive 93/42/EEC.

following the procedure relating to the Annex V.3 of Directive 93/42/EEC, and complying with EN standards of:
EN 980:2008, EN 1041: 2008, EN ISO 14971:2009, EN 60601-1:2005, EN 60601-1-2:2015, EN 1640:2009, EN 15223-2012, ISO11143:2008.

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

Jiangsu Dynamic Medical Technology Co., Ltd

Jiangsu, China, Mar 22, 2020
Place , date

General Manager
Legally binding signature, Function