JOYTECH HEALTHCARE CO., LTD.

EC Declaration of Conformity

Manufacturer:
JOYTECH HEALTHCARE CO. LTD.
Address:No.365, Wuzhou Road, Yuhang Economic Devel
opment Zone, Hangzhou city, 311100 Zhejiang,China

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

We, the manufacturer, herewith declare that the products
Electric Breast pump
Model: LD-202,LD-213,LD-302,LD-305,LD-306,LD-306L
UMDNS-Code: 10485
meet the provisions of Directive 93/42/EEC as amended by Directive 2007/47/EC which apply to them.
The medical device has been assigned to class IIa by rule 11 according to Annex IX of the Directive 93/42/EEC as amended by Directive 2007/47/EC.
It bears the mark

CE 0197

The product concerned has been evaluated under technical files compliance according to Annex VII, and manufactured under a quality management system according to Annex V of Directive 93/42/EEC as amended by Directive 2007/47/EC.

Compliance of the designated product with the Directive 93/42/EEC as amended by Directive 2007/47/EC has been assured via assessment of the quality management system by the Notified Body.

TÜV Rheinland LGA Products GmbH
Tillystraße 2, 90431, Nürnberg, Germany
Certificate No.: DD 60147728 0001
Issue date: 2020-04-07
Expiry date: 2024-05-26

following the procedure relating to the EC Declaration of Conformity set out in Annex V of Directive 93/42/EEC as amended by Directive 2007/47/EC.

This Declaration of conformity is valid in connection with the release document for the respective batch of produced devices.

The above mentioned declaration of conformity is exclusively under the responsibility of
Company: JOYTECH HEALTHCARE CO., LTD.
Address: No.365, Wuzhou Road, Yuhang Economic Development Zone, Hangzhou city, 311100 Zhejiang,China

Hangzhou, July 10, 2020
Place , date

General Manager
Legally binding Signature, Function

JOYTECH HEALTHCARE CO LTD 浙江健拓医疗仪器科技有限公司