No.: MDN-20250711-D001L

Certificate of EU Medical Device Notification

This is to certify that, according to Regulation 2017/745 - medical devices,

Rep Europe GmbH

SRN: DE-AR-000025090

Address: Prinzenallee 7, 40549 Düsseldorf, Germany

has fulfilled all notification responsibility and duty as the European Authorized representative of

Manufacturer: Lingshi International Trade (Hebei) Co., Ltd

SRN: CN-MF-000048514

Address: No.12,Xingda street,Xishiqiao Industrial Zone,Shiqiao Town,Qingyuan District, Baoding City,Hebei Province

The manufacturer has provided with all the appropriate declaration according the Regulation 2017/745 - medical devices requirements including the EC Declaration of Conformity confirming that the medical devices, as stipulated here below, is fulfilling the applicable requirements of Regulation 2017/745.

Where then manufacturer affixes the CE marking to the product listed, they must ensure that all the requirements of the appropriate EU regulation(s) have and continue to be met.

The notification of aforementioned device(s) has been completed by European Authorized representative in Germany, the German Competent Authority has notified the manufacturer's medical device above and has allocated registration. The validity of this document is five years from the date issued.

Product I:	Manual Medical Bed
Registration number:	DE/CA20/00209930
Basic UDI:	697880393LSm2R
Model:	LSm
Product II:	Electric Medical Bed
Registration number:	DE/CA20/00209931
Basic UDI:	697880393LSe29
Model:	LSe
Product III:	Medical Trolley
Registration number:	DE/CA20/00209932
Basic UDI:	697880393LSt37
Model:	LSt

Rep Europe GmbH

11/Jul/2025

Email: info@rep-europe.de

www.rep-europe.de

Digitally signed by Guțan Ghenadie
Date: 2025.09.26 16:17:09 EEST
Reason: MoldSign Signature
Location: Moldova
MOLDOVA EUROPEANĂ

CE Declaration of Conformity

According to Art. 19 of Regulation (EU)2017/745 on Medical Devices

Manufacturer: Lingshi International Trade (Hebei) Co., Ltd

SRN: CN-MF-000048514

Address: No.12,Xingda street,Xishiqiao Industrial Zone,Shiqiao Town, Qingyuan District, Baoding City,Hebei Province, PRC

Email: lingshiguoji123@163.com

Authorized Representative: REP Europe GmbH

SRN: DE-AR-000025090

Address: Prinzenallee 7, 40549 Düsseldorf, Germany

Email: info@rep-europe.de

Standards Applied:

EN ISO 13485: 2016;

EN ISO 14971: 2019;

EN ISO 10993-1:2018;

EN 60601-1:2006/A1:2013 + A2:2021;

EN 60601-2-52:2023

Classification according to MDR Ax. VIII:

Class I., Non-sterile, Non-measuring

Product: Electric Medical Bed

Model: LSe;LSe-1;LSe-2;LSe-3;

LSe-4;LSe-5;LSe-6;LSe-7;LSe-8;

LSe-9;LSe-10;LSe-11;LSe-12;

LSe-13;LSe-14;LSe-15;LSe-16;

Product Type: Medical Device

Basic UDI: 697880393LSe29

Declaration: We, the manufacturer, herewith declare under our sole responsibility that the above-mentioned products meet the provisions of the Regulation (EU) 2017/745 on Medical Devices (MDR). All supporting documentations are retained under the premises of the manufacturer.

Signature: *Tian Jitong*

Name: Jitong Tian

Position: CEO

Signature:

Place and Date of Issue:

Place: Baoding

Date: 11.Jul.2025

CE Declaration of Conformity

According to Art. 19 of Regulation (EU)2017/745 on Medical Devices

Manufacturer: Lingshi International Trade (Hebei) Co., Ltd

SRN: CN-MF-000048514

Address: No.12,Xingda street,Xishiqiao Industrial Zone,Shiqiao Town, Qingyuan District, Baoding City,Hebei Province, PRC

Email: lingshiguoji123@163.com

Authorized Representative: REP Europe GmbH

SRN: DE-AR-000025090

Address: Prinzenallee 7, 40549 Düsseldorf, Germany

Email: info@rep-europe.de

Standards Applied:

EN ISO 13485: 2016;

EN ISO 14971: 2019;

EN ISO 10993-1:2018;

Classification according to MDR Ax. VIII:

Class I., Non-sterile, Non-measuring

Product: Medical Trolley

Model: LSt;LSt-1;LSt-2;LSt-3; LSt-4;LSt-5;LSt-6;LSt-7;LSt-8; LSt-9;LSt-10;LSt-11;LSt-12; LSt-13;LSt-14;LSt-15;LSt-16;

Product Type: Medical Device

Basic UDI: 697880393LSm2R

Declaration: We, the manufacturer, herewith declare under our sole responsibility that the above-mentioned products meet the provisions of the Regulation (EU) 2017/745 on Medical Devices (MDR). All supporting documentations are retained under the premises of the manufacturer.

Signature: Tian Jitong

Name: Jitong Tian

Position: CEO

Signature:

Place and Date of Issue:

Place: Baoding

Date: 11.Jul.2025

CE Declaration of Conformity

According to Art. 19 of Regulation (EU)2017/745 on Medical Devices

Manufacturer: Lingshi International Trade (Hebei) Co., Ltd

SRN: CN-MF-000048514

Address: No.12,Xingda street,Xishiqiao Industrial Zone,Shiqiao Town, Qingyuan District, Baoding City,Hebei Province, PRC

Email: lingshiguoji123@163.com

Authorized Representative: REP Europe GmbH

SRN: DE-AR-000025090

Address: Prinzenallee 7, 40549 Düsseldorf, Germany

Email: info@rep-europe.de

Standards Applied:

EN ISO 13485: 2016;

EN ISO 14971: 2019;

EN ISO 10993-1:2018;

EN 60601-2-52:2023

Classification according to MDR Ax. VIII:

Class I., Non-sterile, Non-measuring

Product: Manual Medical Bed

Model: LSm;LSm-1;LSm-2;LSm-3;
LSm-4;LSm-5;LSm-6;LSm-7;LSm-8;
LSm-9;LSm-10;LSm-11;LSm-12;
LSm-13;LSm-14;LSm-15;LSm-16;

Product Type: Medical Device

Basic UDI: 697880393LSm2R

Declaration: We, the manufacturer, herewith declare under our sole responsibility that the above-mentioned products meet the provisions of the Regulation (EU) 2017/745 on Medical Devices (MDR). All supporting documentations are retained under the premises of the manufacturer.

Signature: *Tian Jitong*

Name: Jitong Tian

Position: CEO

Signature:

Place and Date of Issue:

Place: Baoding

Date: 11.Jul.2025