NINGBO GREETMED MEDICAL INSTRUMENTS CO., LTD.

Declaration of Conformity

Manufacturer:
Ningbo Greetmed Medical Instruments Co., Ltd.
Address: 18F-3, No.1 Building, Wante Business Centre, Hi-Tech Zone, 315042 Ningbo, PEOPLE'S REPUBLIC OF CHINA
Tel: 86-574-87739070 Fax: 86-574-87722360

European Representative:
Shanghai International Holding Corp. GmbH (Europe)
Address: Eiffestrasse 80, 20537 Hamburg, Germany
Tel:+49-40-2513175 Fax:+49-40-255726
Email: shholding@hotmail.com

Digitally signed by Tutunaru Diana
Date: 2021.07.05 11:45:58 MSK
Reason: MoldSign Signature
Location: Moldova

Product Name: Manual Resuscitator
Model: GT012-300A, GT012-300B, GT012-300C
Classification (MDD, Annex IX): Ⅱa (Rule 7 of Annex Ⅸ)
Conformity Assessment Route: Annex V.3

We herewith declare that the above mentioned products meet the transposition into national law, the provisions of the following EC Council Directives and Standards. All supporting documentations are retained under the premises of the manufacturer.

DIRECTIVES

General applicable directives:
Medical Device Directive: COUNCIL DIRECTIVE 93/42/EEC of 14 June 1993 concerning medical devices (MDD 93/42/EEC).
Standard Applied: All applicable harmonized Standard (publish in the Official Journal of the European Communities)
Notified Body: TÜV SÜD Product Service GmbH, Ridlerstr. 65, 80339 MÜnchen, Germany
Identification number: CE0123

(EC) Certificate(s): G20732830046 Rev.01
Expire date of the Certificate: 2024-05-26
Place, Date of Issue: Ningbo , 2020-03-16
Signature:
Name: Li Guirong
Position: General Manager

NINGBO GREETMED MEDICAL INSTRUMENTS CO.,LTD. 宁波格林美康医疗器械有限公司 330291001304