Declaration of Conformity

Manufacturer: RFX + CARE Manufacturing Co., Ltd.
Address: 7 Lanjiang Road, Yuecheng District, Shaoxing, P. R. China

EC-Representative: RFX+Care International A/S

Address: Bakkegaardsvej 408, 3050 Humlebaek, Denmark
Item number: FA200-12570516 / 700905 / 700922 / FA200-12570511 / FA200-12570509 / FA200-12570508 / FA200-12570507 / FA200-12570505 / FA200-12570503 / FA200-12570500
Product Name: Mykit First Aid Kit
UMDNS Code: 11723
Specifications:

Item No.	Name	Specification	Classification
45124	Plaster	56x19mm	I, Rule 4
45125	Plaster	72x25mm	I, Rule 4
45126	Plaster	72x19mm	I, Rule 4
45280	Skin cleansing swab	3x3cm	I Sterile, Rule 1
45191	Dry swab	5x5cm	I Sterile, Rule 4
45107	Elastic bandage	4mx6cm	I, Rule 1
45194	Burn Lint Pad	7x5cm	I Sterile, Rule 4

Classification of Product: Class I sterile, Rule 4
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. RFX + CARE Manufacturing Co., Ltd. is exclusively responsible for the declaration of conformity.

Medical Device Directives

General applicable directives:
Medical Device Directive: COUNCIL DIRECTIVE 93/42/EEC of 14 June 1993 concerning medical devices (MDD 93/42/EEC). Amended by DIRECTIVE 2007/47/EC of 5 September 2007.
Notified Body: TÜV SÜD Product Service GmbH,
Ridlerstrasse 65, 80339 München, Germany
NB Identification number: 0123
(EC) Certificate(s): G2S 067759 0029 Rev.00
Expire date of the Certificate: **2023-11-05**
Start of CE Marking: **2008-11-06**
Place, Date of Issue: **Shaoxing, 2023-06-16**

General Manager (Signature): _____

Date: 2023-06-16