EC Declaration of conformity

Manufacture:

Reliance Medical (Shanghai) Co.,Ltd.
Building 94, Lane 328, Hengyong Road, Jiading District, Shanghai, China.

Whose single Authorized Representative:

Reliance Medical（Ireland）
Unit17, Westlink Industrial Estate, Kylemore Road,Dublin10,Ireland

We, the manufacturer, herewith declare that the p

Plasters

GMDN-Code: 44990

meet the provisions of Directive 93/42/EEC (amended by 2007/47/EC) which apply to them.

The medical device has been assigned to class I sterile according to Annex IX of the Directive 93/42/EEC. It bears the mark

CE 0197

The product concerned has been designed and manufactured under a quality management system according to Annex V of Directive 93/42/EEC (amended by 2007/47/EC).

Compliance of the designated product with the Directive 93/42/EEC(amended by 2007/47/EC) has been assessed and certified by the Notified Body

TÜV Rheinland LGA Products GmbH
Tillystraße 2, 90431, Nürnberg, Germany
Certificate No.: 15051275013
Issue date: 2020-10-09
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(amended by 2007/47/EC).

Application of the abovementioned Annexes and the intervention by the Notified Body is limited to:
the aspects of manufacture concerned with securing and maintaining sterile conditions.

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

Reliance Medical (Shanghai) Co., Ltd

Address: Reliance Medical (Shanghai) Co.,Ltd. Building 94, Lane 328, Hengyong Road, Jiading District, 201806 Shanghai, P.R. China.

Place,date
Shanghai 2021/08/12

Legally binding signature, Function
Jay. Jin. Quality Manager.
< Miaomiao. Jin >