G Declaration of Conformity

Declaration of Conformity V1.0

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CE

Manufacturer:	Shenzhen Shenke Medical Instrument Technical Development Co., Ltd. Bldg 2,5, Mindray Guangming Facility, 1203 Nanhuan Avenue, Yutang Block, Guangming District, 518016, Shenzhen, P.R.China
EC-Representative:	Shanghai International Holding Corp. GmbH (Europe) Eiffestraße 80, 20537 Hamburg, Germany
Product:	Syringe Pump
Model:	BeneFusion SP3, BeneFusion SP3 Ex

We herewith declare that the above mentioned products meet the provisions of the Council Directive 2014/53/EU concerning radio equipment. All supporting documentations are retained under the premises of the manufacturer.

Standards Applied:

☒ EN 60601-1:2006/A1:2013	☒ EN 60601-1-2:2007/AC:2010
☒ EN 62311 :2008	☒ ETSI EN 301 489-1 V2.1.1:2017-02
☒ ETSI EN 301 489-17 V3.1.1:2017-02	☒ EN 300 328 V2.1.1:2016-11
☒ EN60950-1:2006+A11:2009+A1:2010+A12:2011+A2:2013	

Start of CE-Marking:	2017-5-18
Place, Date of Issue:	Shenzhen
Signature:	.....
Name of Authorized Signatory:	Lei Ming
Position Held in Company:	Management Representative

Digitally signed by Alexa Gheorghe
Date: 2023.09.19 09:48:56 EEST
Reason: MoldSign Signature
Location: Moldova