Declaration of Conformity

Name and address of the manufacturer:	Shandong Lianfa Medical Plastic Products Co., Ltd.

No.1 Shuangshan Sanjian Road, 250200, Zhanggiu City, Jinan,

Shandong, PEOPLE'S REPUBLIC OF CHINA

EC Authorized Representative: Linkfar Healthcare GmbH

Niederrheinstraße 71, 40474 Düsseldorf, Germany

We declare under our sole responsibility that :

The medical device: Lancing Device

UMDNS code: 18866

Basic UDI-DI 694951701VA

Trade name

Classification

According to annex VIII(Rule 1) of Regulation EU 2017/745(MDR)

Model: A,B,C,D,E,F,G

SRN Not yet available

We,the manufacture herewith declare that the above-mentioned products meet the provisions of the Regulation EU 2017/745(MDR) . The products meet prospective uses and all supporting documentation is retained under the premise of manufacturer.

Shandong Lianfa Medical Plastic Products Co., Ltd is solely responsibility for the Declaration of Conformity.

Conformity assessment procedure: Annex II & Annex III of EU 2017/745

Declaration of Conformity is valid until: 2021-12-30

Signature:

Name: Yang Lianying Position: President

Place, Date: Jinan, 2021-06

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