山东连发医用塑胶制品有限公司
Shandong Lianfa Medical Plastic Products Co.,Ltd

EU Declaration of Conformity

Manufacturer:	***Whose single Authorized Representative:***
Shandong Lianfa Medical Plastic Products Co., Ltd. No.1 Shuangshan Sanjian Road, Zhangqiu , Jinan City,250200 Shandong P. R. China SRN: CN-MF-000028790	Linkfar Healthcare GmbH Niederrheinstraße 71, 40474 Düsseldorf, Germany SRN: DE-AR-000005107

We, the manufacturer, herewith declare that the products

Product Name:Lancing Devices

Model: LDA; LDC; LDE; LDF; LDC-Ⅱ; LDE-Ⅱ; LDF-Ⅱ

EMDN Code:V9099

Basic UDI-DI: 694951701VA

Intended use:

Assisting the lancet for peripheral blood sampling from the fingertip or the paw for blood glucose testing or other testing utilising a small amount of blood, it is the non-invasive device and does not come into contact with blood, it only makes a brief contact with the skin surface for a few seconds. The lancing device is non-sterile.

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Standards applied:

EN ISO 20417:2021, EN ISO 15223-1:2021,EN ISO 10993-1:2020, EN ISO 10993-5:2009, EN ISO 10993-10:2023, EN ISO 10993-23:2021;EN ISO 14971:2019, EN 62366-1:2015, ASTM F 1980:2016，ASTM D 4169:2016, EN ISO 13485:2016

Meet the provisions of Regulation (EU) 2017/745 which apply to them.

The medical device has been assigned to class I according to Rule 1, Annex VIII of the Regulation (EU) 2017/745. It bears the mark

CE

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

Company: Shandong Lianfa Medical Plastic Products Co., Ltd.

Address: No.1Shuangshan SanjianRoad, Zhangqiu,Jinan City,250200 Shandong P. R. China

Jinan 2025-1-11 — *Place, date*

Lianying Yang|CEO — *Legally binding signature, Function*