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

EC DECLARATION OF CONFORMITY

Name and address of the manufacturer:	**Shandong Lianfa Medical Plastic Products Co., Ltd.** **No.1 Shuangshan Sanjian Road, 250200, Zhangqiu City, Jinan, Shandong, PEOPLE'S REPUBLIC OF CHINA** **SRN: CN-MF-000028790**
EC Authorized Representative:	**Linkfar Healthcare GmbH** **Niederrheinstraße 71, 40474 Düsseldorf, Germany** **SRN: DE-AR-000005107**

We declare under our sole responsibility that

The medical device:	**Sterile Lancet for single use**
UMDNS code:	**10440**
Product code	**01-1330-II, 01-1333-I**
Trade name	/
Classification acc.to MDD Ax.IX:	**Class IIa,rule 6**
Basic UDI-DI	694951700V8

We herewith declare that the above mentioned products meet the provisions of the following EC Council Directives and Standards. All technical documentations are retained under the premises of the manufacturer. We, the manufacturer, are exclusively responsible for this DOC.

Conformity assessment procedure:	**MDD Annex V+Annex VII**
CE Certificate No.:	**G2 036153 0021 Rev.00**
Valid until:	**2024-05-26**
Notified Body:	**TÜV SÜD Product Service GmbH** **Ridlerstraße 65** **80339 MÜNCHEN** **Germany** **CE 0123**

Jinan, 2023-11-04 — Fang Xiao /General Manager

Place, date — Name and function