

Declaration of Conformity

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

Address: No.1 Shuangshan Sanjian Road, 250200, Zhangqiu City, Jinan, Shandong,

PEOPLE'S REPUBLIC OF CHINA

EC Representative: Linkfar Healthcare GmbH

Address: Niederrheinstraße 71, 40474 Düsseldorf, Germany

Products: Lancing device

UMDNS Code:18866

Classification: Class I, Rule 1 According to Annex IX of the MDD

Models and specifications:03-13091

We herewith declare that the above-mentioned products meet the provisions of the following EC Council Directive and Standards (MDD/93/42/EEC). The products meet prospective uses and all supporting documentation is retained under the promise of manufacturer. Our company is exclusively responsible for Declaration of Conformity.

DIRECTIVES

General Applicable Directive:

Medical Device Directive: COUNCIL DIRECTIVE 93/42/EEC concerning medical device (MDD 93/42/EEC)

Notify Body: TÜV SÜD Product Service GmbH, Ridlerstr.65, 80339 Munich, Germany

Identification number: 0123

EC-certificate number: G2 036153 0021 Rev.00

Valid Until: May 05,2024

Signature:_

Name: Fang Xiao

Position:General Manager

Place, Date: Jinan, 2023-11-04