EU Declaration of Conformity

Manufacturer: Ningbo Jenius Polymer Sci-Tech Co., Ltd

No. 2112, Ningheng South Road, Hengxi Town, Yinzhou District, Ningbo, Zhejiang, China (Mainland)
SRN:CN-MF-000026423

whose single Authorized EU-Representative:

MedPath Gmbh
Mies-van-der-Rohe-Strasse 8,80807
Munich,Germany
+49(0)898130 6837
+49(0)89 5485 8884
info@medpath.pro
SRN:DE-AR-000000087

We, the manufacturer, herewith declare that the products:
Tourniquets(JS-TQ001,BASIC UDI-DI:6974491610205)
Tourniquets(JS-TQ003,BASIC UDI-DI:6974491610175)
DESTINATION OF USE:Used to temporarily block venous return during intravenous infusion or blood drawing.

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

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

CE

This Declaration of Conformity covers all medical devices as specified in the product list belonging to this declaration.

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

Ningbo Jenius Polymer Sci-Tech Co., Ltd

Ningbo 2024-09-10
Place, date

Cathy Lee ,General Manger
Legally binding signature, Function

宁波杰尼斯高分子科技有限公司
NINGBO JENIUS POLYMER SCI-TECH CO., LTD