



EU Declaration of Conformity

Doc.No.:FN-QS9.3-RZ-MD-TDC-26070601 Rev.00

Manufacturer:	Finess Healthcare Group Co.,Ltd 6th Floor, Building C, Jiaotong BOBO International, No.221 Yongfeng West Road, Haishu District, Ningbo, 315000, China	SRN No.:	CN-MF-000004124
European Representative:	SUNGO Europe B.V. Fascinatio Boulevard 522, Unit 1.7, 2909V A Capelle aan den IJssel, The Netherlands	SRN No.:	NL-AR-000000247
Brand Holder	Tendercare		
Product Name:	Razor	Brand Trade Name	DISPOSABLE RAZOR
Product REF:	FN-RZ010105	Brand REF	430746
Basic UDI-DI:	697391169RZ01RH		

Classification (MDR, Annex VIII): Class I Non sterile,Rule 1
Conformity Assessment Route: Article 19+Annex II+Annex III

We herewith declare that this EU Declaration of Conformity is issued under the sole responsibility of the manufacturer.The products above are in conformity with the Medical Device Regulation (EU) 2017/745.All supporting documentation is retained under the premises of the manufacturer.

General Applicable Regulation:

Medical Device Regulation (EU) 2017/745 and other applicable EU standards and MDCG Guidance.

Place,Date of Issue: Ningbo,2026-07-06

Signature:

Finess Healthcare Group Co.,Ltd
Quality/Regulatory Department:

Name: Ruby Fu

Ruby Fu

Position: PRRC