



## FINESS HEALTHCARE GROUP CO.,LTD

### EU Declaration of Conformity

Doc.No.:FN-QS9.3-WTD-MD-TDC-26070604 Rev.00

|                          |                                                                                                                                                                 |                  |                               |
|--------------------------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------|------------------|-------------------------------|
| Manufacturer:            | <b>Finess Healthcare Group Co.,Ltd</b><br>6th Floor, Building C, Jiaotong BOBO International, No.221 Yongfeng West Road, Haishu District, Ningbo, 315000, China | SRN No.:         | CN-MF-000004124               |
| European Representative: | <b>SUNGO Europe B.V.</b><br>Fascinatio Boulevard 522, Unit 1.7, 2909V A Capelle aan den IJssel, The Netherlands                                                 | SRN No.:         | NL-AR-000000247               |
| Brand Holder             | Tendercare                                                                                                                                                      |                  |                               |
| Product Name:            | Wooden Tongue Depressor                                                                                                                                         | Brand Trade Name | WOODEN TONGUE SPATULA<br>15CM |
| Product REF:             | FN-WTD010101                                                                                                                                                    | Brand REF        | 388993                        |
| Basic UDI-DI:            | 697391169WTD01TB                                                                                                                                                |                  |                               |

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

Name: Ruby Fu

Quality/Regulatory Department:

Position: PRRC

Ruby Fu