# Declaration of Conformity

## 1. Products

### Fingerbob

**Intended purpose:**

It's suitable for securing primary wound dressing, perfect for all types of finger injury.

**Basic UDI-DI:** 694109403889FY

| General Device Group | Product name | Product code | Description-ENG | Rule | Risk Class |
|---|---|---|---|---|---|
| (Non-sterile) Bandage and Tape product | Finger bob | WIN9044000 | ASKINA FINGER BOB WHITE | Rule 1 | Class I non-sterile |
| | | WIN9044019 | ASKINA FINGER BOB COLOURED | | |
| | | WIN9044027 | ASKINA FINGER BOB WHITE | | |
| | | WIN9044035 | ASKINA FINGER BOB COLOURED | | |
| | | 507220 | FINGER BOB STD. WHITE/TUBE. FUMOUZE | | |
| | | 507219 | FINGER BOB STD. COLOUR/TUBE. FUMOUZE | | |
| | | WINMX1049 | FINGER BOB STD. WHITE/TUBE. BOMEDYS | | |
| | | WINMX1050 | FINGER BOB STD. COLOUR/TUBE. BOMEDYS | | |
| | | WIN1016B | JLB FINGER BOB - STD. BLUE | | |
| | | WIN1016W | JLB FINGER BOB - STD. WHITE | | |
| | | WIN1017B | JLB FINGER BOB - LARGE BLUE | | |
| | | WINMX1009 | FINGER BOB LARGE BLUE BULK MARIAX | | |
| | | WINMX1040 | FINGER BOB STD. WHITE/TUBE. MARIAX | | |
| | | WINMX1043 | FINGER BOB STD. WHITE/TUBE. GENERICOS | | |
| | | WINMX1044A | FINGER BOB STD. COLOR/TUBE. HILEFARMA | | |
| | | WINMX1044 | FINGER BOB STD. WHITE/TUBE. HILEFARMA | | |
| | | WIN1017W | JLB FINGER BOB - LARGE WHITE | | |
| | | WIN9044159 | ASKINA FINGER BOB LARGE | | |
| | | WIN9044540 | FINGER BOB MEDIUM. WHITE- BULK | | |
| | | WIN9044558 | FINGER BOB LARGE WHITE(IN BULK) | | |

## 2. Manufacturer

Name: Winner Medical Co., Ltd.
Address: Winner Industrial Park, No.660 Bulong Road, Longhua District, 518109 Shenzhen, CHINA
SRN Number：CN-MF-000005692

## 3. Authorised Representative

Name: Shanghai International Holding Corp, GmbH (Europe)
Address: Eiffestraße 80, 20537 Hamburg, Germany
SRN Number: DE-AR-000000001

## 4. Statement

We hereby declare that the above mentioned products with CE marking meet the provisions of Regulation (EU) 2017/745 of the European Parliament and of the Council. The CE marking is subject to the general principles set out in Article 30 of Regulation (EC) No 765/2008.The Declaration of conformity is issued under the sole responsibility of the manufacturer.

## 5. Conformity Assessment Procedure

EU declaration of conformity (Article 19) + technical documentation (Annexes II and III) of Regulation (EU) 2017/745 concerning medical device for Class I (Non-sterile) medical device.

**For and on behalf of Winner Medical Co.,Ltd.**

Xiaomeng Yang

**Xiaomeng, Yang**
**Management Representative**
**Shenzhen, China.**
**Issued Date (YY-MM-DD):2021.10.08**