

Doc. No.	KSW/TD-FBN-017	Title	EU Declaration of Conformity of Finger Bob		
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EU Declaration of Conformity

Manufacturer Name: Kingstar Industries (Wuhan) Co., Ltd.

Manufacturer Address: No.5 Xiaoxiangxi Road, Jinyinhu, Dongxihu District, Wuhan, Hubei, China, 430040

Location of Manufacturer: Wuhan City, Hubei Province, China.

Authorized Representative: Shanghai International Holding Corp. GmbH (Europe)

Address of their Registered Place of Business: Eiffestrasse 80, D-20537, Hamburg, Germany

Location be established: Germany

Basic UDI-DI code: 6973554891431000WJ **Registered Trade Name / Mark:** Kingstar

Name of the device: Finger Bob

CND Code: M03030103, TUBULAR NET BANDAGES

UMDNS Code: 10291, Bandages, Tubular

Intended Purpose: Intended to be externally worn on a finger/limb to provide support or local pressure to a part of the body, especially a joint, for various preventative/therapeutic applications (e.g., support soft tissue injuries) while enabling movement.

Risk Class of the Device: Class I, based on Rule 1 of ANNEX VIII of Regulation (EU) 2017/745.

The Conformity Assessment Procedure Performed: According to Article 19 of the Regulation (EU) 2017/745, draw up this EU declaration of conformity which contain the information set out in Annex IV of the Regulation (EU) 2017/745.

CS used or Standard applied: N/A

Identification of the device: Please find in Annex I

Declaration: This declaration of conformity is issued under the sole responsibility Kingstar Industries (Wuhan) Co., Ltd. We hereby declare that the medical device specified above meet the provision of the Regulation (EU) 2017/745 for medical device. This declaration is supported by the quality system approval to ISO 13485 by TÜV SÜD Product Service GmbH

All supporting documentation is retained at the premises of the manufacturer.

Signed for and on behalf of:

Place of Issue: Wuhan City, Hubei Province, China.

Date of Issue: 2020-02-28 Print Name: Aigun Zhou

Function: Management Representative

Signature:



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Annex I --- Identification of the Device Covered by the EU Declaration of Conformity

1. Identification of the Device

Table --- Identification of the Device

No.	WERO REF	Kingstar Article Number	Specification	Packaging Configuration
1	110019	5142470010	finger bob, blue, 1.5 g/pc, Stretched Length: 55cm, Width: 7cm; Diameter: 2.5cm	40 pcs/PE bag, 125 bags/case
2	110025	/	finger bob, blue, 1.5 g/pc, Stretched Length: 55cm, Width: 7cm; Diameter: 2.5cm	5 Stk./Pkg
3	110027	/	finger bob, blue, 1.5 g/pc, Stretched Length: 55cm, Width: 7cm; Diameter: 2.5cm	40 Stk/Pkg

2 Photograph of Finger Bob



Photo 1 --- Finger Bob in PE bag



Photo 2 --- Finger Bob