

# EU DECLARATION OF CONFORMITY



manufacturer: ZARYS International Group sp. z o.o. sp.k.  
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SRN: PL-MF-000000410

We declare under our sole responsibility that a medical device:

**yellowTAPE**  
**kinesiotape**

sizes\*: 2,5 cm x 5 m, 5 cm x 5 m, 7,5 cm x 5 m, 5 cm x 32 m

(\*detailed list of products covered by this declaration is available in document TD-29-I.1.1.b-1– Identification – Annex 1, batch code - certificate of analysis of production batch DZDO-01– Annex 2)

classification:

- **class I, rule 1** (in accordance with Annex VIII of Regulation (EU) 2017/745)

Basic UDI-DI: **59079968M0599QB**

intended purpose: Device intended for taping on the surface of the body in accordance with the principles of kinesiotaping technique (dynamic plastering) as a prevention of injuries or alleviation of diseases of the musculoskeletal system.

is in conformity with Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices.

The device described above meets all applicable provisions of the Annex I of Regulation (EU) 2017/745. Conformity assessment procedure has been performed in accordance with Article 52 (7).

The medical device covered by the present declaration of conformity complies with European standards. The list of supervised standards is included in document TD-29-I.4.c-1 - Annex 3.

place and date of issue: Zabrze, 27.09.2023  
name: Wioletta Wójcik  
position: Product Manager

PRODUCT MANAGER  
ZARYS International Group sp. z o.o. sp.k.

Handwritten signature of Wioletta Wójcik in blue ink.  
Wioletta Wójcik

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signature  
(on behalf of the President of the General Partner's  
Management Board)

