

## EC Declaration of Conformity

*Manufacturer:*

K-JUMP HEALTH CO., LTD  
NO.56, Wu Kung 5th Rd., New Taipei Industrial Park,  
New Taipei City 24890, Taiwan

*whose single Authorized Representative:*

POLYGREEN GERMANY GmbH  
Ruhlsdorfer Straße 95, D-14532, Stahnsdorf,  
Germany

**DIMDI Code:** DE/0000044710

We, the manufacturer, herewith declare that the products

KP-6027 (SC 6027, SC 6027A), KP-6260 (SC 6200), KP-6262 (SC 6262), KP-6241 (SC 6400), KP-6750 (SC 6750), KP-6860 (SC 6800), KP-7012 (SC 7012), KP-7160 (SC 7100), KP-7130 (SC 7130), KP-7160 (SC 7160), KP-7030 (SC 7300), KP-7020 (SC 7400 NFC), KP-7530 (SC 7530), KP-7660 (SC 7660, SC 7600), KP-7620 (SC 7620), KP-7670 (SC 7670), KP-7590 (SC 7701), KP-7920 (SC 2100), KP-6020 (SC 8100), KP-6520 (SC 8900),

### NON-INVASIVE ELECTRONIC SPHYGMOMANOMETER

**UMDNS-Code: 16-173**

meet the provisions of Directive 93/42/EEC which apply to them.

The medical device has been assigned to class IIa according to Annex IX of the Directive 93/42/EEC. It bears the mark

**CE 0197**

The product concerned has been designed and manufactured under a quality management system according to Annex II (without the Annex II.4) of Directive 93/42/EEC and the essential requirement of Annex I pertaining to medical devices.

Compliance of the designated product with the Directive 93/42/EEC has been assessed and certified by the Notified Body

**TÜV Rheinland LGA Products GmbH**  
**Tillystraße 2, 90431 Nürnberg, Germany**

Registration No.: HD 60148105 0001

Issue date: 2020-07-28

Expiry date: 2024-05-26

following the procedure relating to the EC Declaration of Conformity set out in Annex II and Annex VII of Directive 93/42/EEC.



This Declaration of conformity is valid in connection with the release document for the respective batch of produced devices.

The above mentioned declaration of conformity is exclusively under the responsibility of

## K-JUMP HEALTH CO., LTD

K-JUMP HEALTH CO.,LTD.

*Tiffany Su*

Place, Date

Taipei, 2022-09-01

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*Tiffany Su/Sales Manager*

