Übereinstimmungserklärung

Statement of Conformity



Hiermit wird bestätigt, dass die Produkte der

It is hereby confirmed that the products of

Ossenberg GmbH Kanalstraße 79 48432 Rheine Deutschland

identisch mit denen der von

are identical with those of

HEBEI Healthplus Medical Device Co., Ltd.

No. 1, Chuangye Street, Southwest Industrial District,
Matou Ecological Industry Park, Handan City
Hebei, 056046
China

gelieferten Artikel sind.

deliveres articles.

| Artikel-Nr. | Produktbezeichnung | Artikel-Nr. | Produktbezeichnung |
|----------------|---|-------------|---|
| HEBEI | HEBEI Healthplus Medical Device Co., Ltd. | Ossenberg | Ossenberg GmbH |
| Healthplus | | GmbH | |
| Medical Device | | | |
| Co., Ltd. | | | |
| article no. | Product name | article no. | Product name |
| HEBEI | HEBEI Healthplus Medical Device Co., Ltd. | Ossenberg | Ossenberg GmbH |
| Healthplus | | GmbH | |
| Medical Device | | | |
| Co., Ltd. | | | |
| 16si | Vierfußgehhilfe | 16si | Vierfuß-Gehhilfe in silber mit weichem |
| | | | Schaumstoffgriff – 100 kg |
| | Quad Cane | | Quadruped walking aid in silver with soft |
| | | | foam grip – 100 kg |

Rheine, den 09.01.2024

Ort, Datum / place, date

Dr. Thomas Schreder

Name und Unterschrift Verantwortliche Person / Name and signature PRRC



EU Declaration of Conformity

The following description for the medical device,

| Device information | Description | | |
|-------------------------------------|---|--|--|
| Registered trade name and address | HEBEI Healthplus Medical Device Co., Ltd. No. 1, Chuangye Street, Southwest Industrial District, Matou Ecological Industry Park, Handan City, Hebei, 056046, China Tel:+86-0310-2111888 | | |
| Authorized representative | Y. Sung Handelsvertretung Toulouser Allee 9, 40211 Duesseldorf, Germany | | |
| Common device name | Canes | | |
| Product and trade name | Healthplus User Friendly 訊普 成 Medical Products | | |
| UMDNS code | 10560, Canes | | |
| GMDN code | 31110: Walking Aid | | |
| | 17105: Walking Aid Handgrip | | |
| Single Registration Number (SRN) | Manufacturer: CN-MF-000013062 / Authorized representative: DE-AR-000005142 | | |
| Basic UDI-DI | 697322200MXCANEHK | | |
| Risk class of the device | Class I | | |
| Common Specification (CS) | Canes, Folding Seat Canes, Height Adjustable Canes, | | |
| references | Quad Canes, Offset Quad Canes, Walking Aids, | | |
| | Walking Aids with Handgrip, Walking Sticks | | |
| Intended purpose | An all- or partially-wheeled mobility aid in the form of a | | |
| (GMDN definition) | waist-high framework with handgrips and four or three legs | | |
| | designed to provide a stable support to a person with a disability, | | |
| | a geriatric or an infirm person when standing or ambulating; it | | |
| | may also be used during rehabilitation. It is constructed of | | |
| | lightweight metal, plastic, and rubber materials in the shape of a | | |
| | mobile support structure which the user holds and pushes in from | | |
| | of them as they walk. It is non-foldable and includes a braking | | |
| | mechanism; it may be width-adjustable and some types may | | |
| | include a seat on which the user can rest. | | |
| Conformity assessment procedure | Quality Management System | | |
| performed and identification of the | ISO 13485:2016 by NQA | | |
| certificates issued by notified | ISO 9001:2015 by NQA | | |
| body, if applicable | | | |
| Name and identification number of | ISO 13485, NQA Certificate Number: 35429 | | |
| the notified body, if applicable | ISO 9001, NQA Certificate Number: 46296 | | |

Issued date: March 5, 2020

Version: V1.0



Regulation 2017/745/EU and, if applicable, with any other relevant Union legislation that provides for the issuing of this EU declaration of conformity. The device is in conformity with conformity assessment procedure for Class I devices that should be carried out, as a general rule, under the sole responsibility of manufacturers in view of the low level of vulnerability associated with such devices. Thus, manufacturers of class I devices, other than custom-made or investigational devices, shall declare the conformity of their products by issuing an EU declaration of conformity referred to in Article 19 "EC declaration of conformity" after drawing up the technical documentation set out in Annexes II and III of the Regulation.

For the evaluation regarding Class I device (Risk class in accordance with the Rule 1 set out in Annex VIII of the **Regulation**), the following harmonized standards are applied:

- EN ISO 14971:2019 Medical devices Application of Risk Management
- EN ISO 15223-1:2021 Medical device Symbols to be used with medical device labels, labeling and information to be supplied Part 1: General requirements
 - EN 62366-1:2015 Medical devices Part 1-- Application of usability engineering to medical devices

The following Union authorized representative is stated to the declaration:

Y. Sung Handelsvertretung

Toulouser Allee 9, 40211 Duesseldorf, Germany

(Company name / Registered place of business)

The following person is exclusively responsible for the compliance of declaration:

HEBEI Healthplus Medical Device Co., Ltd.

No. 1, Chuangye Street, Southwest Industrial District, Matou Ecological Industry Park,

Handan city, Hebei, 056046, China

(Manufacturer's name/ Registered address)

Dickson Su / General Manager

(Name/Function)

(Legal Signature)

March 5, 2023

(Date of issue)

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