

EU DECLARATION OF CONFORMITY

1. Basic UDI-DI: **2090425RollzFlex2K2**
2. Item codes: 3010RF0010, 3010RF0012, 3010RF0013, 3010RF0014, 3011RF0010.
3. Name: Rollz International B.V.
Adress: Rotterdamseweg 402M
2629 HH Delft
The Netherlands
4. This declaration of conformity is issued under the sole responsibility of the manufacturer (or installer):
Rollz International B.V.
Rotterdamseweg 402M
2629 HH Delft
The Netherlands
SRN: NL-MF-000002978
5. Rollz Flex rollator.



6. The object of the declaration described in point 5 is a **class 1** medical device and is in conformity with **REGULATION (EU) 2017/745 OF THE EURO PEAN PARLIAMENT AND OF THE COUNCIL of 5 April 2017 on medical devices.**

7. References to the relevant harmonised standards used, or references to the specifications in relation to which conformity is declared:

- **NEN-EN 11199-2:2021 Walking aids manipulated by both hands – Requirements and test methods - Part 2: Rollators.**
- **NEN-EN 12182:2012 Assistive product for people with disability – General requirements and test methods.**
- **NEN-EN 1985:1998 Walking aids – General requirements and test methods.**
- **NEN-EN 14971:2019 Medical devices – Application of risk management to medical devices.**

8. Certified by SGS Taiwan Ltd.

Accreditation No: 1053

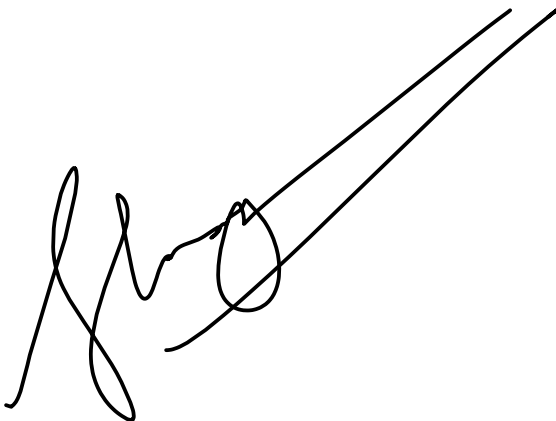
Certificate No: **HQ60047/2021 (ISO 11199-2)**

9. Signed for and on behalf of

Rollz International B.V.

Delft, 11-10-2021

Martijn Schaaper, CEO Rollz International

A handwritten signature in black ink, appearing to be 'M. Schaaper', is written over a diagonal line that extends from the bottom left towards the top right of the page.