EU Declaration of Conformity

according to the Regulation (EU) 2017/745 on Medical Devices (MDR) Annex IV

Manufacturer Name:	LIAM DEVELOPMENTS BV
Manufacturer Address:	JACHTWAGEN 29 3897 AM ZEEWOLDE (NL)
SRN (Single Registration Number):	NL-MF-000032442
Name of Device:	**Rollbuddy® Indoor Rollator**
BASIC UDI / GMN:	8720726413RB013J
Product Code (SKU)	RB01ALU / RB01CPR / RB01BLK
Risk Classification:	Class I according to rule 1 set out in annex VII of Regulation (EU) 2017/745
Conformity Assessment Route:	Manufacturer uses the procedures for the CE labeling of their products according to the Regulation MDR 2017/745.

This Declaration of Conformity is issued under the sole responsibility of LIAM DEVELOPMENTS BV. We hereby declare that the medical device(s) specified above meet the provision of the Regulations (EU) MDR 2017/745 for medical devices. All supporting documentation is retained at the premises of the manufacturer.

Signature:

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Menno Vaartjes
(Director)

Place and date of issue:
Zeewolde, 06-01-2023