Manufacturer´s Declaration of Conformity

CE marking in accordance with the Medical Device Regulation (EU) 2017/745

Manufacturer´s name: **Manufacturer's Adress:**	Lopital Nederland B.V. Laarakkerweg 9, 5061 JR, Oisterwijk The Netherlands
Manufacturer's SRN (Single Registration Number):	NL-MF-000004372
Brand Name:	Lopital
	
Medical device:	
Model number(s):	Lotus, Lotus Compact & Lotus XL
Device Description:	59008200, 59008250 & 59008270
Basic UDI-DI:	Active Standup Aid 8720256103615590082000PZ, 8720256103325590082500NL & 8720256103646590082700U2
Classification:	Class I
Conforms to regulation:	Medical Device Regulation (EU) 2017/745
Standards applied:	NEN-EN-ISO 14971:2019
The product has been tested and evaluated in accordance with the following standards:	ISO 10535:2021 (corrigendum 2023-11) IEC 62366-1:2015 + A1:2020
The product has been designed and manufactured under a certified quality management system in accordance with:	UNI-CEI-EN-ISO 13485:2016

This declaration of conformity is issued under the sole responsibility of Lopital. We hereby declare that the medical device(s) specified above meet the provision of the Regulation (EU) MDR 2017/745 for medical devices.

Signature:	Date:	Place:
	18-05-2026	Oisterwijk
Patrycja Wojciuk, Manager quality & Regulatory	dd-mm-yyyy	

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