

Manufacturer's Declaration of Conformity

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

Lopital Nederland B.V.

Manufacturer's Adress:	Laarakkerweg 9, 5061 JR, Oisterwijk The Netherlands	
Manufacturer's SRN (Single Registration Number):	NL-MF-000004372	
Brand Name:	Lopital	
Medical device: Model number(s): Device Description: Basic UDI-DI:	Nemo 51005100 Mobile tilting Shower-Toiletchair 872025610304251005100DP	
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 17966:2016 EN-IEC 62366-1:2015	
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:
	12-12-2024	Oisterwijk —
Jan Van Megen, CEO	dd-mm-yyyy	

Document Number: DOC5100-2024-02

Manufacturer's name: