

Manufacturer's Declaration of Conformity

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

Manufacturer's name:	Lopital Nederland B.V.
Manufacturer's Address:	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:	Timo 54006020, 54006021, 54006025, 54006030, 54006034 & 54006035 Mobile Shower-Toiletchair 872025610307354006020JY, 872025610308054006021HS, 872025610309754006025PY, 872025610310354006030DP, 872025610347954006034TH & 872025610311054006035CR
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 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:


Jan Van Megen, CEO

Date:
12-12-2024

dd-mm-yyyy

Place:
Oisterwijk
