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 Adress:	Laarakkerweg 9, 5061 JR, Oisterwijk The Netherlands
Manufacturer's SRN (Single Registration Number):	NL-MF-000004372
Brand Name:	Lopital
	
Medical device:	Sirocco basic & Sirocco deluxe
Model number(s):	61002500 & 61002510
Device Description:	Wall mounted Shower Stretcher with electrically powered high/ low adjustment
Basic UDI-DI:	8720256103561610025 00L7 & 8720256103578610025 10S8
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 60601-1-2 (2015)+A1(2021) NEN-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: DOC2500-2024-03