

DECLARATION OF CONFORMITY

We are solely responsible for declaring that the Medical Devices mentioned in this statement are of Low-Risk Class (Class I) and comply with the requirements of the European Regulation 2017/745 and where appropriate, the standards and legislation referred to.

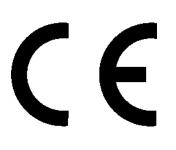
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COMPETENT AUTHORITY:	National Organization for Medicines	
CONTACT DETAILS:	Address: Mesogeion 284, PS 15562, Cholargos Phone: 2132040000 Website: https://www.eof.gr/	

LIST OF PRODUCTS COVERED BY THIS DECLARATION					
PRODUCT	CODE	BASIC UDI-DI	INTENDED USE	RULE	
TELESCOPIC RAMP 7FT	0805400	521300690ramps17LV	USE BY PEOPLE WITH MOBILITY PROBLEMS	1	
ROLL UP RAMP	0805401	521300690ramps17LV	USE BY PEOPLE WITH MOBILITY PROBLEMS	1	
SUITCASE RAMP 3FT (WITH ANTI- SLIPERY TAPE) 91CM	0805402	521300690ramps17LV	USE BY PEOPLE WITH MOBILITY PROBLEMS	1	
SUITCASE RAMP 5FT (WITH ANTI- SLIPERY TAPE) 152 CM	0805403	521300690ramps17LV	USE BY PEOPLE WITH MOBILITY PROBLEMS	1	

CONFORMITY ASSESSMENT PROCEDURE		
According to Annexes II & III of Regulation (EU) 2017/745		

APPLIED STANDARDS & LEGAL REQUIREMENTS		
ISO 13485:2016, ISO 9001:2015, EN ISO 15223-1:2021, (EU) 2017/745		



FOR APPROVAL		
NAME:	SVOURAKI MARIA	
POSITION:	CEO	
PLACE:	CHANIA	
DATE:	24/05/2021	
SIGN:	DIA.	