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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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

CE

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