

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
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LIST OF PRODUCTS COVERED BY THIS DECLARATION

PRODUCT	CODE	BASIC UDI-DI	INTENDED USE	RULE
OVER BED TABLE RECLINING (BEIGE)	0809243	521300690TABLES219V	CABINET NEAR THE BED & TABLE BED, FOR THE FACILITY OF THE PATIENT	1
OVER BED TABLE FIXED (BEIGE)	0808234	521300690TABLES219V	CABINET NEAR THE BED & TABLE BED, FOR THE FACILITY OF THE PATIENT	1
OVER BED RECLINING TABLE WHEELED	0806645	521300690TABLES219V	CABINET NEAR THE BED & TABLE BED, FOR THE FACILITY OF THE PATIENT	1
OVER BED FIXED TABLE WHEELED	0806646	521300690TABLES219V	CABINET NEAR THE BED & TABLE BED, FOR THE FACILITY OF THE PATIENT	1
OVER BED TABLE FOR HOSPITAL USE (MAX 23KG)	0811377	521300690TABLES219V	CABINET NEAR THE BED & TABLE BED, FOR THE FACILITY OF THE PATIENT	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 20417:2021, ISO 14971:2019, ISO 15223-1:2021, (EU) 2017/745



FOR APPROVAL

NAME:	SVOURAKI MARIA
POSITION:	CEO
PLACE:	CHANIA
DATE:	23/11/2022
SIGN:	

