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
BEDSIDE CABINET, WOODEN FOR HOSPITAL USE	0805620	521300690bedsides15FE	SUITABLE FOR HOSPITAL USE	1
BEDSIDE CABINET (WOOD STICKER) FOR HOSPITAL USE	0805621	521300690bedsides15FE	SUITABLE FOR HOSPITAL USE	1
BEDSIDE CABINET, WOODEN, WITH HEIGHT ADJ. TABLE FOR HOSPITAL USE	0805623	521300690bedsides15FE	SUITABLE FOR HOSPITAL USE	1
BEDSIDE CABINET	0805900	521300690bedsides15FE	SUITABLE FOR HOSPITAL USE	1

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

APPLIED STANDARDS & LEGAL REQUIREMENTS
ISO 13485:2016, ISO 9001:2015, ISO 14971:2019, EN 62366:2015, ISO 10993-1:2018, EN ISO 15223-1:2021, (EU) 2017/745

CE

FOR APPROVAL	
NAME:	**SVOURAKI MARIA**
POSITION:	**CEO**
PLACE:	**CHANIA**
DATE:	**21/12/2024**
SIGN: