

## EU DECLARATION OF CONFORMITY

<b>Manufacturer name</b>	Optium Medikal Ltd.Sti.			
<b>Registered address</b>	Körkün Mah. Hidayet Cad. No:23 OĞUZELİ GAZİANTEP TURKEY			
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<b>E-mail</b>	<a href="mailto:info@optium.com.tr">info@optium.com.tr</a>			
<b>Brand</b>				
<b>Directive</b>	Regulation (EU) 2017/745 on medical devices			
<b>Conformity Assessment</b>	Regulation (EU) 2017/745 on medical devices EK-IV EU DECLARATION OF CONFORMITY (EK II & III)			
<b>Classification of the product as the medical device:</b>	According To Annex VIII of Regulation (EU) 2017/745 on medical devices Class I Other (nonsterile, without measuring function)			
<b>Basic UDI-DI</b>	868432544044OTH2Z			
<b>Products</b>	<b>Model</b>	<b>Product Name</b>	<b>UDI-DI Number</b>	<b>EMDN Code</b>
	<b>MR 39</b>	Medical Records Storage Cabinet	8684325441650	V0880
	<b>LC 15</b>	Linen Cabinet	8684325441704	V0880
	<b>MHT 32</b>	Filling Cabinet	8684325441742	V0880
	<b>CI 19</b>	Medicine Cabinet	8684325441810	V0880
	<b>CM 35</b>	Medicine and Instrument Cabinet	8684325441827	V0880
	<b>CM 37</b>	Instrument and Medicine Cabinet	8684325441834	V0880
	<b>CM 38</b>	Medical Cabinet	8684325441841	V0880
<b>Description and function designation:</b>	Metal cabinet, including all applicable accessories.			

WE HEREWITH DECLARE THAT THOSE ABOVE PRODUCTS WITH CE MARKING WHICH ARE MANUFACTURED BY OUR COMPANY ALL COMPLY WITH REGULATION (EU) 2017/745 ON MEDICAL DEVICES, AND REALIZE THEIR EXPECTED USES. ALL CE FILES HAVE BEEN CERTIFIED BY THE COMPANY, CONSEQUENTLY THEIR AUTHENTICITY HAS BEEN QUARANTEED.

### Harmonised Standarts

The said products fulfills the requirements of these harmonized technical standards which were used for assessing of conformity

A statement that the declaration of conformity is issued under the responsibility of the manufacturer.

Place, Date of Issue : GAZİANTEP / 12.08.2021  
Approved By : Ahmet DAL / General Manager  
Stamp and Signature :

