


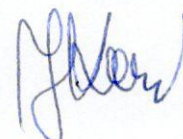
EU DECLARATION OF CONFORMITY

1	Registered trade name and the address of the registered place of business.	
	JJW Spółka Jawna J. Karpiński, J. Rudnicki, W. Lewin Plac Zgody 6 05-520 Konstancin-Jeziorna, Poland	
2	A statement that the EU declaration of conformity is issued under the sole responsibility of the manufacturer.	
	We hereby state that the EU declaration of conformity is issued under the sole responsibility of the JJW Spółka Jawna J. Karpiński, J. Rudnicki, W. Lewin	
3	The Basic UDI-DI as referred to in Part C of Annex VI.	
	Generic name of the product: Prophylactic medical socks	
	Models:	
	Trade name of the model	Basic UDI-DI
	DeoMed Cotton Silver	5901050DMCottonSilverRW
	DeoMed Bamboo	5901050DMBambooCQ
	Smart SeaCell	5901050SmartSeaCell9J
	Medic Deo Cotton	5901050MDCottonNS
	Medic Deo Silverwool	5901050MDSilverwoolE6
DeoMed Wool	5901050DMWool36	
4	Product and trade name, product code, catalogue number or other unambiguous reference allowing identification and traceability of the device covered by the EU declaration of conformity, such as a photograph, where appropriate, as well as its intended purpose. Except for the product or trade name, the information allowing identification and traceability may be provided by the Basic UDI-DI referred to in point 3.	
	Generic name of the product: Prophylactic medical socks	
	Models:	
	Trade name of the model	Model reference code
	DeoMed Cotton Silver	DMS
	DeoMed Bamboo	DMB
	Smart SeaCell	SSZ
	Medic Deo Cotton	MDC
Medic Deo Silverwool	MSW	
DeoMed Wool	DMW	
5	Risk class of the device in accordance with the rules set out in Annex VIII.	
	Class I Rule 1	
6	A statement that the device that is covered by the present declaration is in conformity with this Regulation and, if applicable, with any other relevant Union legislation that provides for the issuing of an EU declaration of conformity.	
	The device models to which the declaration relates comply with REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 5 April 2017 on medical devices, amending Directive 2001/83/EC, Regulation (EC) No 178/2002 and Regulation (EC) No 1223/2009 and repealing Council Directives 90/385/EEC and 93/42/EEC (Text with EEA relevance).	

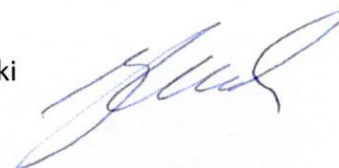


7	<p>References to any CS used and in relation to which conformity is declared.</p> <p>EN ISO 15223-1: 2016 [PN-EN ISO 15223-1: 2017-02] Medical devices - Symbols for use on medical device labels, markings and information provided with them - Part 1: General requirements</p> <p>EN 1041: 2008 + A1: 2013 [PN-EN 1041 + A1: 2013-12] Information provided by the manufacturer of medical devices</p> <p>EN ISO 14971: 2019 [PN-EN ISO 14971: 2020-05] Medical devices. Application of risk management to medical devices.</p> <p>and with other documents:</p> <p>ZN-JJW-W-013: 2019 DEOMED COTTON SILVER socks - manufacturing process</p> <p>ZN-JJW-W-005: 2011 DEOMED BAMBOO socks - manufacturing process</p> <p>ZN-JJW-W-009: 2015 SMART SEACELL socks - manufacturing process</p> <p>ZN-JJW-W-001: 2018 MEDIC DEO COTTON socks - manufacturing process</p> <p>ZN-JJW-W-010: 2015 MEDIC DEO SILVERWOOL socks - manufacturing process</p> <p>ZN-JJW-W-011: 2017 DEOMED WOOL socks - manufacturing process</p>
8	<p>Where applicable, the name and identification number of the notified body, a description of the conformity assessment procedure performed and identification of the certificate or certificates issued.</p> <p>Not applicable</p>
9	<p>Where applicable, additional information.</p> <p>Not applicable</p>
10	<p>Place and date of issue of the declaration, name and function of the person who signed it as well as an indication for, and on behalf of whom, that person signed, signature.</p>

Janusz Karpiński
co-owner



Jerzy Rudnicki
co-owner



Włodzimierz Lewin
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Konstancin-Jeziorna, on April 30, 2021

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Place, date

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Name, function and signature