



## EC Declaration of Conformity

<b>Manufacturer</b>	Name	MGsolutions Inc.
	Address	1F,5F, 10, Donggyo-ro 23-gil, Mapo-gu, Seoul, Republic of Korea
	SRN	KR-MF-000051033
<b>European Representative</b>	Name	CMC Medical Devices & Drugs S.L.
	Address	C/ Horacio Lengo n18, C.P 29006, Málaga-Spain
	SRN	ES-AR-000000293
<b>Sole Responsibility Statement</b>	This EU Declaration of Conformity is issued under the sole responsibility of the manufacturer, MG solutions Inc,	
<b>BASIC UDI-DI</b>	8800332570026	
<b>Product Identification</b>	Product Name	Moti Physio Med
	Intended Use	Moti Physio Med captures and visualizes a person's static and dynamic posture and joint positions. It produces 3D representations and surface-contour images to illustrate body alignment and before/after changes. The qualitative or semi-quantitative indicators generated serve as reference information to support physical therapist and other medical professionals when observing musculoskeletal function and designing exercise or rehabilitation programs. The product neither delivers calibrated measurements nor makes diagnoses and treatment decisions. All clinical interpretation and decisions remain entirely with the professional user.
<b>Classification</b>	Class I, according to Rule 11 of Annex VIII of Regulation (EU) 2017/745.	
<b>Conformity Statement</b>	The device that is covered by this declaration is in conformity with Regulation (EU) 2017/745 on medical devices.	
<b>Common Specifications</b>	Not applicable	
<b>Notified Body</b>	Not applicable. (The conformity assessment procedure for Class I devices does not require the involvement of a Notified Body.)	
<b>Additional Information</b>	None	

※ All supporting documentation is retained under the premises of the manufacturer and manufacturer is exclusively responsible for the declaration of conformity.

Signature: \_\_\_\_\_ Jae-hyun Park, 2025-12-16  
President of MGsolutions Inc.