





All products will be accompanied by the following statement in line with Annex XIII of (EU) 2017/745

DMO Custom-Made Medical Device
Statement by Manufacturer

This custom-made medical device has been manufactured to the prescription attached and is intended for the exclusive use of:

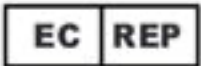
		
		
		
		

The device conforms to the essential requirements (Part II of the UK MDR 2002, Annex I [as modified by part of schedule 2A to the UK MDR 2002])

The device conforms to the GSPR requirements of Annex I of MDR (Medical Devices Regulation) (2017/745) (MDR)



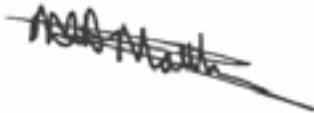
DM Orthotics Ltd, Unit 2, Cardrew Way, Cardrew industrial Estate, Redruth, Cornwall, TR15 1SS, UK



Emergo Europe, Westervoortsedijk 60, 6827 AT Arnhem, The Netherlands

Approved By: Martin Matthews, PRRC Date:22/03/2023

Signed:



Year of Manufacture: 2023

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