

**EC** – Declaration of Conformity

Manufacturers Name:	Orfit Industries N.V.		• •		•••	
					• •	
SRN (Single Registration	BE-MF-000007872		• •		• •	
Number):			• •		•••	
Manufacturers Address:	Vosveld 9a, 2110 Wijnegem, Belgium				• •	
Manufacturers Address.	Vosvela Ja, ZIIO Wijnegeni, Deigiani		• •		• •	
					•••	
Basic UDI-DI:	5420028700414Y					
			• •	•	• •	
Name of the Device(s):	Raycast <sup>®</sup> High Precision - Accessories Head Supports - Low Density for	push-pin	• •		• •	0
	hardware					
					• •	
			• •	•	• •	1
Intended use:	Patient Positioning and Immobilization for Radiation Oncology		• •	•	• •	
Product code(s):	32414, 32415, 32412, 32413, 32410, 32411, 32409					
			• •	•	• •	1
			• •	•	• •	1
Classification:	Class I, according the rules of Annex VIII					
Conformity assessment route: Orfit Industries N.V. uses the following procedures for the CE-labelling of their						
	products according the Regulation MDR 2017/745:			•	•••	
	products according the negatiation MDN 2017/145.					
			• •	•	• •	
	Class I: EC conformity declaration according to Annex IV.	· · 0	• •	•	• •	1
					••	
Applied norms:	ISO 13485:2016				• •	
	ISO 14971:2019	0	• •	•	• •	1
	ISO 15223-1:2021	÷ 0	• •	•	• •	1
	ISO 10993-5:2009				• •	1
	ISO 10993-10:2010				• •	
				•	• •	1

This declaration of conformity is issued under the sole responsibility of Orfit Industries N.V. We hereby declare that the medical device(s) specified above meet the provisions of the Regulation (EU) MDR 2017/745 for medical devices. This declaration is supported by the Quality System approval ISO 13485:2016 issued LRQA. All supporting documentation is retained at the premises of the manufacturer.

Eddy Marivoet, Wijnegem, 22 May 2024

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Quality Assurance & Regulatory Affairs Manager

