

## **EC** – Declaration of Conformity

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Manufacturers Name:	Orfit Industries N.V.	· •			
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SRN (Single Registration	BE-MF-000007872	• 0	• •	٠	٠
Number):			• •	٠	۰
Number j.					1
					÷.
Manufacturers Address:	Vosveld 9a, 2110 Wijnegem, Belgium				
					•
Basic UDI-DI:	5420028700465A	· · 0	• •	•	۰
	5 1200207 00 1007 (	0	• •	•	۰
Nouse of the Device (a)	Developt <sup>®</sup> Lliph Duppinian Association Dupped Lloyd Components	• •			
Name of the Device(s):	Raycast <sup>®</sup> High Precision Accessories Prone Head Supports				ī.
Intended use:	Patient Positioning and Immobilization for Radiation Oncology		• •	•	•
intended dse.			• •	٠	•
		· · · 0 0	• •	٠	۰
Product code(s):	32393, 32380, 32379		• •		
					÷.
Classification:	Class I, according the rules of Annex VIII				•
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Conformity assessment route	: Orfit Industries N.V. uses the following procedures for the CE-labellin	g of their	• •		•
comorning assessment routes		gormen			Ĵ.
	products according the Regulation MDR 2017/745:				
	Class I: EC conformity declaration according to Annex IV.		• •	٠	
	class I. Le comorning declaration according to Annex IV.		• •	٠	٠
			• •		•
Applied norms:	ISO 13485:2016				1
	ISO 14971:2019				÷.
	ISO 15223-1:2021	0		•	•
		• •	• •	•	•
	ISO 10993-5:2009		• •	٠	•
	ISO 10993-10:2010		• •		
			• •	1	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 by LRQA. All supporting documentation is retained at the premises of the manufacturer.

Eddy Marivoet, Wijnegem, 21 May 2024

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Quality Assurance & Regulatory Affairs Manager

