Leader in thermoplastic innovations

EC – Declaration of Conformity

Manufacturers Name:	Orfit Industries N.V.
SRN (Single Registration Number):	BE-MF-000007872
Manufacturers Address:	Vosveld 9a, 2110 Wijnegem, Belgium
Basic UDI-DI:	5420028700244Y
Name of the Device(s):	HP PRO® SOLUTION
Intended use:	Patient Positioning and Immobilization for Radiation Oncology

Product code(s): 25000/1, 25000/22, 25000/23, 25000/25, 25000/20, 25000/30, 25000/18, 25000/26, 25000/1/MR, 25000/27, 25000/16, 25000/36, 25000/33, 25000/16/MR, 25000/6, 25000/58/20, 25000/39/20, 25000/17/A, 25000/17/B, 25070/10, 25071/10, 25000/90, 25000/91, 25000/92, 25000/94, 25100/16MI+N, 25100/16Mi+N/NH, 25113/24MI+N, 25105/2MI+N, 25105/2MI+N/NH, 25115/24MI+N, 25115/24MI+N/NH, 25116/24MI+N, 25103/16MI+N/NH, 25101/16MI+N, 25101/16MI+N/NH, 25110/24MI+N, 25107/2MI+N/NH, 25106/2MI+N, 25106/2MI+N/NH, 25112/24MI+N, 25000/75, 25000/27/MR, 25000/76, 25000/62, 25000/67, 25000/69, 25000/69/A, 25000/69/B, 25000/7, 25000/7/20, 32138, 25100/2MI+/NH, 25100/2MI+, 25105/2MI+, 25105/2MI+/NH, 25000/26, 25000/70, 25000/71, 25000/68, 25000/61, 25000/83, 25000/84, 25000/88**,** 25000/81, 25000/89, 25000/50, 25000/82, 25000/87, 25000/34, 25000/38, 25000/53, 25000/43, 25000/63, 25000/77, 25000/78, 25000/79, 25000/80**,** 25000/8, 25000/64, 25000/40 25113/24MI+N/NH, 25117/24MI+N, 25118/24MI+N, 25100/2MI+N/NH, 25109/24MI+N, 25111/24MI+N, 28002

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:

Class I: EC conformity declaration according to Annex IV.

Applied norms:

ISO 13485:2016
ISO 14971:2019
ISO 14971:2019/A11:2021
ISO 15223-1:2021
ISO 10993-1:2025
ISO 10993-5:2009
ISO 10993-10:2021

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, 8 September 2026

Quality Assurance & Regulatory Affairs Manager

ORFIT INDUSTRIES
Vosveld 9A | B-2110 Wijnegem | Belgium
T (+32) (0)3 326 20 26
welcome@orfit.com