260320/ha-prs-1

EU DECLARATION OF CONFORMITY

Manufacturer:	Oy Klippan Ab Santaradantie 8 FI-01370 Vantaa SRN: FI-MF-000052644
Manufactured for:	High Adventure Oy Pulttitie 14 00880 HELSINKI
Basic UDI-DI:	641764002900AM
Name of Device:	HARRT Pro PRS – Restraint System
Product code:	HA-PRS
Intended Use:	For use in conjunction with basket stretchers
Classification:	Class 1, according to annex VIII, of MDR 2017/745

Conformity Claim

The device/accessory covered by this declaration is in conformity with Regulation (EU) 2017/745.

- EN ISO 14971:2019 + A11:2021
- EN ISO 15223-1

We hereby declare that this EU Declaration of Conformity is issued under the sole responsibility of the manufacturer, and the medical device specified above meet the provision of the Regulation (EU) MDR 2017/745 for medical devices.

Oy Klippan Ab or representative will make available upon request all applicable technical documentation to allow assessment of conformity of its products

Vantaa 20.3.2026

Oy Klippan Ab

Hans Bäckström
CEO

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