CE

Declaration of Conformity

Manufacturer name:	Danrehab A/S
Manufacturer address:	Vejlevej 15, Ølholm DK-7160 Tørring
Manufacturer SRN:	DK- 66261417
Device Name:	FISK
Device Description:	Therapeutic stool

Part Number:	24726
Basic UDI:	571506418090401GH
HMI No.:	27726
Classification:	N/A
Conforms to regulation:	GPSD (General Product Safety Directive).
The Product has been tested, and evaluated in accordance with the following standards:	GPSD (General Product Safety Directive).
The product has been designed and manufactured under a certified quality management system in accordance with:	EN ISO 13485:2016

We hereby declare that the medical device defined above complies with the GPSD (General Product Safety Directive). Classification has been confirmed with the Danish Medicines agency.

Maria W Laursen

Head of Quality and Compliance, MDR
27-10-2025