EU declaration of conformity

The devices covered by this declaration are in conformity with Regulation (EU) 2017/745 on medical devices.

Device name	Protac® MyRest
Basic UDI-DI	5714821007403M
EMDN	Y99
Intended purpose	The body pillow is used to stimulate the different senses in children and adults. The various sensory stimuli can be influenced via applying weight and pressure to the sense of touch (the tactile sense) in the skin and the sense of movement (the proprioceptive sense) in muscles and joints. The body pillow is also intended as alleviation for restless legs.
Risk class	Class I, rule 1

This EU declaration of conformity is issued under the sole responsibility of the manufacturer.

Manufacturer	Protac A/S Niels Bohrs Vej 31D DK-8660 Skanderborg www.protac.dk
SRN	DK-MF-000000963
Place	Stilling, Denmark
Date of issue	24-09-2024
Name and function	Trine Danielsen, QA/RA manager

Signature on behalf of Protac A/S