EC Declaration of Conformity

Manufacturer:
KayserBetten GmbH & Co KG
Rieper Str. 12
D-29683 Bad Fallingbostel
Germany
SRN: DE-MF-000006910

We declare under our sole responsibility that the products listed below comply with the relevant provisions of the following regulations.

Product designation	**Risk class** (EU) 2017/745 Annex VIII	**Basis UDI-DI**
KayserBetten TIMMY 1	I	426038961TIMMY1HL
KayserBetten TIMMY 2	I	426038961TIMMY2HN
KayserBetten TIMMY easyLift	I	426038961TIMMYEASYLIFT62
KayserBetten TIMMY Telescope	I	426038961TIMMYTELESKOP9S

Purpose:
Children's care beds are used for chronically ill and/or physically/mentally impaired children and adolescents for nursing, therapeutic care and as sleeping and reclining places.

Regulations:
(EU) 2017/745 Medical Devices Regulation

Conformity assessment procedure (EU) 2017/745:
Annex II, Annex III, Annex IV, Annex V

Applied applicable parts of the standards:
BS EN 50637:2018
BS EN 60601-1:2013
BS EN 60601-2-52:2016
BS EN 716-1:2019

In the event of relevant changes to the above-mentioned medical devices, this Declaration of Conformity loses its validity in accordance with the requirements of EK-Med Decision 3.9 A4 of the ZLG.

Bad Fallingbostel, 23.06.2022

Torsten Kappenberg / Managing Director

Jens Lübben / Head of Medical Technology, PRRC

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