

EU overensstemmelseserklæring / EU-Declaration of Conformity

i henhold til artikel 19 i forordning (EU) 2017/745 om medicinsk udstyr
according to article 19 of medical device regulation (EU) 2017/745

Firmanavn / Company name:

Wulff MED TEC GmbH

SRN:

DE-MF-000010006

Vi erklærer under eget ansvar, at det/de produkt(er), der er anført i nedenstående tabel, og som Wulff Med Tec GmbH først markedsførte, opfylder de gældende bestemmelser i forordning (EU) 2017/745 om medicinsk udstyr.

Madrasser til hospitalssenge giver patienterne øget hvile- og sovekomfort. Madrasserne kan understøtte behandlingen og forebyggelsen af liggesår.

Produktet/produkterne er blevet identificeret som medicinsk udstyr i klasse I i henhold til regel 1 i bilag VIII til forordning (EU) 2017/745 om medicinsk udstyr.

Adresse / Address:

Hennstedter Straße 3
25779 Fedderingen
Tyskland / Germany

We declare under sole responsibility, that the product/s listed in the chart below which was/were first placed on the market by Wulff Med Tec GmbH meet/s the applicable provisions of the medical-device-regulation (EU) 2017/745.

Mattresses for hospital beds are used to improve the resting and sleeping comfort of patients. The mattresses can support the therapy and the prevention of pressure injuries.

The product/s was/were identified as a medical device in risk class I according to rule 1 in annex VIII of the medical-device-regulation (EU) 2017/745.

**Grundlæggende UDI-DI
Basic UDI-DI**

40508244PWRUG

**Produktnavn
Product name**

MediMatt 30

**Artikelnr.
Article no.**

4PWR

Gyldighedsperiode / Validity period:

12.02.2025 – 11.02.2027


Melchior Meyer (Feb 12, 2025 07:26 GMT+1)
Melchior C. Meyer
(Managing Director)
(Administrerende direktør)



Fedderingen, 12.02.2025

MediMatt EU Declaration of Coformity_DA

Final Audit Report

2025-02-12

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