

In compliance with Article 52 No. 7 of the EU-Regulation EU 2017/745
of the Council for Medical Products (of April 5th, 2017)

SEGUFIX[®]-Bandagen,
Das Humane System GmbH & Co. KG
(SRN: DE-MF-000027702),

hereby declares bindingly and under own responsibility
that the fundamental requirements according to Annexes I of
the regulation are met for the following medical product:

Item 2205, Item 2305, Item 2405, Item 2105, Item 2005
SEGUFIX[®]-Foot Restraint

in the sizes and versions offered according to the 2024 price list.
Basic-UDI-DI = PP01531220529 / risk class = I

Intended use:

- gradually adjustable restraint from loose to tight
of one or both feet in bed
- physically restraining patients

The conformity assessment procedure according to
Annexes II and III of the EU regulation EU 2017/745
is applied. This declaration is valid from the
issuing date until May 24th, 2026.

Jesteburg, May 25th, 2024

Date of issue



R. W. de Sánchez
– Executive Management –

SEGUFIX[®]-Bandagen • Das Humane System GmbH & Co. KG

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of the Council for Medical Products (of April 5th, 2017)

SEGUFIX[®]-Bandagen,
Das Humane System GmbH & Co. KG
(SRN: DE-MF-000027702),

hereby declares bindingly and under own responsibility
that the fundamental requirements according to Annexes I of
the regulation are met for the following medical product:

Item 2705, Item 2805, Item 2905
SEGUFIX[®]-Foot Restraint Velcro[®]

in the sizes and versions offered according to the 2024 price list.
Basic-UDI-DI = PP01531270535 / risk class = I

Intended use:

- gradual positioning from loose to tight of one or both feet in bed
- not a restraint (note: some jurisdictions consider Velcro[®] products
as a restraint, should the patient not be able to remove them
without help)

The conformity assessment procedure according to
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