

EU Declaration of Conformity

Herewith we declare under our sole responsibility that the following Class I medical device(s) (acc. Rule 1, Annex VIII) comply with the Regulation (EU) 2017/745.

Based on the conformity assessment, the declaration of conformity was issued according to Annex IV of Regulation (EU) 2017/745. Due to risk class 1 and according to Article 52 (7) of Regulation (EU) 2017/745 the manufacturer is entitled to conduct the conformity assessment procedure independently. An evaluation by a notified body is not required.

The implemented Quality Management System fulfills the requirements of EN ISO 13485:2016.

Spinomed[®]

Basic UDI-DI: 4026398Y060309034G

SRN of the Manufacturer: DE-MF-000007092

Intended use: Spinomed® is a brace designed to actively

relieve load and correct the lumbar spine /

thoracic spine in the sagittal plane.

Common Specifications: Not applicable.

We expressly state that the "registered"-sign "®" is not part of the name and is only used to identify a registered trademark, that is why it may appear at different positions.

Bayreuth, 10.01.2024

Stefan Weihermüller, PRRC medi GmbH & Co. KG

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This declaration is valid until: 10.01.2027.