

**Konformitätserklärung
Declaration of Conformity**

Wir

We

**B. Braun Melsungen AG
Carl-Braun-Str. 1
34212 Melsungen
Deutschland/Germany
SRN DE-MF-000000201**erklären in eigener Verantwortung,
dass das Produkt / die Produkte**Perifix® LOR
Perifix® LOR NRFit®**Spritze für die Loss-of-Resistance (LOR)-
Methode in der Regionalanästhesie
(Artikelnummern und Basic UDI-DI siehe Anlage I)mit den Anforderungen der Medizinprodukte
Verordnung (EU) 2017/745 übereinstimmen**Konformitätsbewertungsverfahren
nach Anhang IX****Klassifizierung**
gemäß Anhang VIII der oben genannten
Verordnung
Klasse I steril**Benannte Stelle**
TÜV SÜD Product Service GmbH
Kennnummer 0123**Gültig bis 2026-12-14**
gemäß gültigem EU-Zertifikat
(Nr. G11 012974 0626)hereby declare in our sole responsibility
that the product/s**Perifix® LOR
Perifix® LOR NRFit®**Loss of Resistance (LOR) device for use in
Regional Anesthesia
(article numbers and Basic UDI-DI see attachment I)are in conformity with the requirements of the
Medical Device Regulation (EU) 2017/745**Conformity Assessment Procedure
according to annex IX****Classification**
according to annex VIII of the Regulation named
above
Class I sterile**Notified Body**
TÜV SÜD Product Service GmbH
Identification number 0123**Valid until 2026-12-14**
according to our valid EU Certificate
(No. G11 012974 0626)

Anlage I / Attachment I**Basic UDI-DI: 403923900000238936**

Art.-Nr. / Art. No.	Produktname / Product name	Klasse / Class
4637100	Perifix® LOR	Is (sterile)
4637110	Perifix® LOR NRFit®	Is (sterile)
4638107	Perifix® LOR	Is (sterile)
4638110	Perifix® LOR NRFit®	Is (sterile)

Document amendment information

Version	Description of the changes
1.0	Initial Version under 2017/745 MDR

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This document is signed electronically in compliance with the B. Braun electronic signature policies and procedures by following persons:

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