

We,

**BSN medical GmbH
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Germany
(SRN: DE-MF-000005787)**

hereby declare under our sole responsibility, that this product family complies with the applicable requirements of the regulation (EU) 2017/745 of the European parliament and of the council on medical devices (MDR).

Product Name:

Leukotape® K

Basic UDI-DI:

4042809400351848X

Intended purpose:

**Leukotape® K is indicated for usage in kinesiology-type
taping techniques. Application fields include:**
- Lymphedema
- Muscular pain
- Disordered tonicity

**The device is for single use only.
Intended Users are Health Care Professionals and
patients. Use of device is not restricted to specific
populations.**

Conformity assessment route: **Annex II+III**

Classification rule: **1**

Classification: **I**

The Declaration of Conformity is performed in accordance with the quality management system according to EN ISO 13485, and fulfils the general safety and performance requirements of the regulation (EU) 2017/745.

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Martin Spengler
Director Regulatory Affairs Hamburg
BSN medical GmbH



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