

We,

**BSN medical GmbH  
Schützenstr. 1-3  
22761 Hamburg  
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:

**Leukoplast® skin sensitive**

Basic UDI-DI:

**4042809400351178G**

Intended purpose:

**Leukoplast® skin sensitive is intended for fixation of wound dressings and additional securement of devices, such as catheters, tubes, probes or drainages. It is particularly suitable for highly sensitive, delicate, cracked, fragile, elderly, infant, cortisone/medication-damaged, diabetic skin and for use on all body parts, including frequently moved or highly contoured parts of the body.**

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.

Declaration of conformity issued: 05.02.2025

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Compiled and released:

Hamburg, 05.02.2025  
Martin Spengler  
Director Regulatory Affairs Hamburg  
BSN medical GmbH



Article	Description	REF
76173-00000-02	LEUKOPLAST SKIN SENSITIVE PLASTIC SNAPRING 1.25CMX2.6M 1 DA NL EN FI FR DE EL IT NO PT ES SV 24/BOX	76173-00
76173-00001-02	LEUKOPLAST SKIN SENSITIVE PLASTIC SNAPRING 2.5CMX2.6M 1 DA NL EN FI FR DE EL IT NO PT ES SV 12/BOX	76173-01
76173-00006-02	LEUKOPLAST SKIN SENSITIVE PLASTIC SNAPRING 2.5CMX1M 1 DA NL EN FI FR DE EL IT NO PT ES SV 12/BOX	76173-06
76175-00000-02	LEUKOPLAST SKIN SENSITIVE PLASTIC SPOOL 2.5CMX2.6M 12 DA NL EN FI FR DE EL IT NO PT ES SV 12/BOX	76175-00