

Declaration of Conformity MDR LP Compress Absorbent Soft NST INT

JBX.40049822.01 Page 1 of 2

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® compress absorbent soft

Basic UDI-DI:

404280940049822**A**K

Intended purpose:

Leukoplast® compress absorbent soft is intended for skin and wound cleansing, absorption of secretion/exudates. As mechanical barrier it is suitable as a primary or secondary dressing in order to cover a wound and to provide cushioning. The device is for short-term use.

Conformity assessment route: Annex II+III

Classification rule:

Classification:

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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Compiled and released:

Hamburg, 09.03.2023 Martin Spengler
Director Regulatory Affairs Hamburg
BSN medical GmbH

Madir Mula



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JBX.40049822.01 Page 2 of 2

Article	Description	REF
71281-00000-00	LEUKOPLAST COMPRESS ABSORBENT SOFT NON STERILE 10CX10CM WHITE 100X1 DA NL EN FI FR DE ID IT NO PT ES SV	71281-00
71281-00001-00	LEUKOPLAST COMPRESS ABSORBENT SOFT NON STERILE 10CX20CM WHITE 100X1 DA NL EN FI FR DE ID IT NO PT ES SV	71281-01
71281-00002-00	LEUKOPLAST COMPRESS ABSORBENT SOFT NON STERILE 15CX25CM WHITE 100X1 DA NL EN FI FR DE ID IT NO PT ES SV	71281-02
71281-00003-00	LEUKOPLAST COMPRESS ABSORBENT SOFT NON STERILE 20CX20CM WHITE 100X1 DA NL EN FI FR DE ID IT NO PT ES SV	71281-03
71281-00004-00	LEUKOPLAST COMPRESS ABSORBENT SOFT NON STERILE 20CX40CM WHITE 50X1 DA NL EN FI FR DE ID IT NO PT ES SV	71281-04