

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:

404280940049821AH

Intended purpose:

The device 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.

Conformity assessment route: **Annex IX, Chapter I**

Classification rule:

4

Classification:

Is

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.

This declaration is only valid in conjunction with the current E.C. certificates issued by DEKRA Certification GmbH (Id. No. 0124), Handwerkstraße 15, 70565 Stuttgart, Germany.

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Hamburg, 15.02.2023
Martin Spengler
Director Regulatory Affairs Hamburg
BSN medical GmbH



Article	Description	REF
71280-00000-00	LEUKOPLAST COMPRESS ABSORBENT SOFT STERILE 10CX10CM WHITE 25X1 NL EN FI FR DE ID IT NO PT ES SV	71280-00
71280-00001-00	LEUKOPLAST COMPRESS ABSORBENT SOFT STERILE 10CX20CM WHITE 25X1 NL EN FI FR DE ID IT NO PT ES SV	71280-01
71280-00002-00	LEUKOPLAST COMPRESS ABSORBENT SOFT STERILE 15CX25CM WHITE 15X1 NL EN FI FR DE ID IT NO PT ES SV	71280-02
71280-00003-00	LEUKOPLAST COMPRESS ABSORBENT SOFT STERILE 20CX20CM WHITE 25X1 NL EN FI FR DE ID IT NO PT ES SV	71280-03
71280-00004-00	LEUKOPLAST COMPRESS ABSORBENT SOFT STERILE 20CX40CM WHITE 12X1 NL EN FI FR DE ID IT NO PT ES SV	71280-04