

Declaration of Conformity Leukoplast® soft white

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We,

BSN medical GmbH Quickbornstrasse 24 20253 Hamburg Germany (SRN: not issued yet)

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

Product name: Basic UDI-DI:

Leukoplast® soft white (not defined yet)

Intended purpose:

Refer to clinical evaluation

Annex II+III

Conformity assessment route: 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.

Certificate No 50708-14-00 (EN ISO 13485)

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Compiled and released:

Hamburg, 16.01.2020 Martin Spengler Director Regulatory Affairs Hamburg BSN medical GmbH

Havi Reh



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Article	Description	REF	
76450-00026-00	LEUKOPLAST SOFT WHITE 19X40MM 500 DE DA EN FI FR IT NL NO PT SV ES INJECTION DRESSING	76450-26	