

## **Declaration of Conformity** Leukoplast® swab ball gauze

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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® swab ball gauze

Basic UDI-DI: 404280940048588AT

Intended purpose: Leukoplast® swab ball gauze 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 IX, Chapter I Classification rule: 4

Classification: ls

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.

Date of Issue: 12.05.2023

Compiled and released:

Hamburg, 12.05.2023 Martin Spengler Director Regulatory Affairs Hamburg BSN medical GmbH

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