

Declaration of Conformity Leukomed® I.V. nonwoven

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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:

Leukomed® I.V. non-woven

Basic UDI-DI: 4042809400416208D

Intended purpose: Leukomed® I.V. non-woven is intended for fixation of I.V.

catheters and covering and protection of puncture sites.

Conformity assessment route: Annex IX, Chapter I

Classification rule: İs 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. 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.

Number of E.C. certificate: 50708-60-04

Declaration of conformity issued: 28.03.2024

Compiled and released:

Hamburg, 28.03.2024 Martin Spengler Director Regulatory Affairs Hamburg

BSN medicăl GmbH

Gallin Mula



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Article	Description	REF
72389-00000-04	LEUKOMED INTRAVENOUS STERILE 6X8CM WHITE 50 NL EN DE IT JA ES HOSP.	72389-00