Declaration of Conformity Leukotape® P

We.

BSN medical (Pty) Ltd 30 Gillits Road: 3610 Pinetown **South Africa** (SRN: ZA-MF-000022653)

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:

Leukotape® P

Basic UDI-DI:

4042809400356709D

Intended purpose:

Leukotape® P inelastic tapes are intended for functional taping as treatment and prevention of injuries to muscles, ligaments and joints, rigid joint immobilization and sports

injuries (strains and sprains.

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.

European Authorized Representative:

BSN medical GmbH Schützenstr. 1-3 22761 Hamburg

Germany (SRN: DE-AR-000005791)

Declaration of conformity issued: 30.05.2023

Compiled and released:

Pinetown, 30.05.2023

Julie Maharaj

BSN medical (Pty) Ltd



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Article	Description	REF
76168-00000-05	LEUKOTAPE 3.8CMX13.7M TAN 1 EN DE FR ES PT IT NL SV NO DA FI ZH AR P MICRO SERRATED	76168-00
76169-00000-05	LEUKOTAPE 3.8CMX13.7M TAN 1 EN DE FR ES PT IT NL SV NO DA FI ZH AR P BIG PACK (6/BOX) MICRO SERRATED	76169-00