We,

BSN medical GmbH
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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:	**Uniflex®Universal by Actimove®**
Basic UDI-DI:	**404280940039689AW**
Intended purpose:	**Uniflex® Universal by Actimove® is intended to provide light support or compression as well as fixation and dressing retention in post-operative or post-traumatic usage. Application fields include:** **• Light support in the treatment of minor soft tissue and joint injuries (e.g. sprains, strains)** **• Light compression (e.g. in edema management to reduce effusion and swelling)** **• Secure dressings (e.g. when treating wounds) or fixation (e.g. of casts or splints)**
Conformity assessment route:	**Annex II+III**
Classification rule:	**1**
Classification:	**I**

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.

Declaration of conformity issued: 13.03.2025

Compiled and released:

Hamburg, 13.03.2025
Martin Spengler
Director Regulatory Affairs Hamburg
BSN medical GmbH

Article	Description	REF
02017-00000-06	UNIFLEX UNIVERSAL BY ACTIMOVE 6CMX5M WHITE 1 EN DE FR ES PT IT NL SV NO DA FI ZH AR 10/BOX	02017-00
02018-00000-06	UNIFLEX UNIVERSAL BY ACTIMOVE 8CMX5M WHITE 1 EN DE FR ES PT IT NL SV NO DA FI ZH AR 10/BOX	02018-00
02019-00000-06	UNIFLEX UNIVERSAL BY ACTIMOVE 10CMX5M WHITE 1 EN DE FR ES PT IT NL SV NO DA FI ZH AR 10/BOX	02019-00
02020-00000-06	UNIFLEX UNIVERSAL BY ACTIMOVE 12CM X 5M WHITE 1 EN DE FR ES PT IT NL SV NO DA FI ZH AR (10/BOX)	02020-00
02021-00000-06	UNIFLEX UNIVERSAL BY ACTIMOVE 15CMX5M WHITE 1 EN DE FR ES PT IT NL SV NO DA FI ZH AR (5/FOIL BAG)	02021-00