

We,

**BSN medical GmbH
Quickbornstr. 24
20253 Hamburg
Germany
(SRN: DE-MF-000005787)**

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:

Tensospray®

Basic UDI-DI:

4042809400376119B

Intended purpose:

**The device is a multiple use item to provide an adhesive film on the skin. Application fields include:
- fixation of tapes / bandages**

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.

Date of Issue: 04.05.2021

Compiled and released:

Hamburg, 04.05.2021
Martin Spengler
Director Regulatory Affairs Hamburg
BSN medical GmbH





Declaration of Conformity
Tensospray®

JBX.40037611.03
Page 2 of 2

Article	Description	REF
71602-00000-04	TENSOSPRAY 300 ML 1 NL EN FR DE IT ES	71602-00