

# Declaration of conformity for medical device

Our company,

Ofa Bamberg GmbH  
Laubanger 20  
D - 96052 Bamberg

declares hereby as responsible manufacturer and under sole responsibility that the medical devices listed in Annex comply with all relevant requirements of Regulation (EU) 2017/745 (MDR) in its current consolidated form.

A notified body is not involved in the conformity assessment procedure.

<b>Classification:</b>	I
(according to Annex VIII, Rule 1)	
<b>Single Registration Number (SRN):</b>	DE-MF-000008470
<b>Basic-UDI-DI:</b>	4018839K029J
<b>Conformity assessment:</b>	according to Article 52 & Annex II - IV
<b>CE- labelling:</b>	CE
<b>Validity of the declaration of conformity:</b>	August 31, 2030
<b>Intended purpose:</b>	

Medical compression stockings for the treatment of lymphological diseases, lipedema and scar therapy.

The validity of this declaration of conformity ends with a new declaration of conformity.

Ofa Bamberg GmbH  
Bamberg, 02.02.2026



Stephan Börner  
Managing Director



Dr. Fabian Bohnen  
Managing Director &  
Person responsible for regulatory compliance  
according to article 15, MDR

## Annex

Product	REF-number (1.-4. or 1.-6. digit)
Belsana® Classic Extra CCL I	2224
Belsana® Impuls CCL I	2225
Belsana® Classic Extra CCL II	2244
Belsana® Impuls CCL II	2245
Belsana® Classic Extra CCL III	2264
Belsana® Impuls CCL III	2265
Belsana® Impuls CCL IV	2275
Belsana® Impuls CCL II	2745
Belsana® Impuls CCL III	2765