Spring® medical

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 consolidated form.

Classification:	I
(according to Annex VIII, Rule 1)	
Single Registration Number (SRN):	not yet available
Basic-UDI-DI:	4018839K019G
Conformity assessment:	according to Article 52 & Annex II - IV
CE- labelling:	CE
Validity of the declaration of conformity:	May 25, 2025

We also confirm the compliance with the following laws, regulations and quality standards: MDR, MPDG, RAL-GZ 387/1+2. The validity of this declaration of conformity ends with a new declaration of conformity.

Ofa Bamberg GmbH
Bamberg, 26.05.2021

Rainer Kliewe
Managing director

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

Spring® medical

Annex

Product	REF-number (1.-4. or 1.-6. digit)
Spring® deluxe fino Kkl 1	7181
Spring® deluxe fino Kkl 2	7182
Spring® deluxe fino Kkl 1Maß	7281
Spring® deluxe fino Kkl 2 Maß	7282
Spring® deluxe classico Kkl 1	7152
Spring® deluxe cassico Kkl 2	7154
Spring® deluxe classico Kkl 1 Maß	7252
Spring® deluxe classico Kkl 2 Maß	7254
Spring® deluxe vigento Kkl 2	7134
Spring® deluxe vigento Kkl 2 Maß	7234
Spring® deluxe vigento Kkl 3 Maß	7235