# 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.

| | |
|---|---|
| **Classification:** | I |
| (according to Annex VIII, Rule 1) | |
| **Single Registration Number (SRN):** | DE-MF-000008470 |
| **Basic-UDI-DI:** | 4018839K019G |
| **Conformity assessment:** | according to Article 52 & Annex II - IV |
| **CE- labelling:** | CE |
| **Validity of the declaration of conformity:** | July 01, 2025 |

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

Ofa Bamberg GmbH
Bamberg, 01.11.2021

Rainer Kliewe
Managing director

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

## 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 |