# 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:** | 4018839K049N |
| **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, 22.11.2021

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) |
|---|---|
| Compresso Fix (14-22 mmHg) | 0370 |
| Compresso Fix (23-32 mmHg) | 0372 |