

Declaration of Conformity

Product Designation: Slings, Arm

UMDNS- Code: 13-622

Model No's: 3087, 3187, 3289, 4089, 4189

Issue Date: Oct. 12, 2018 Valid Date: Oct. 12, 2022

We herewith declare that the products listed above are in compliance with the requirements set out in the Council Directive on the harmonization of the Laws of the Member States concerning Medical Device Directive 93/42/EEC.

Conformity Assessment Procedure: MDD 93/42/EEC Annex VII

Classification of the Product: Class I Rule: MDD 93/42/EEC Annex IX Rule 1

Applied Harmonized Standards: EN ISO9001:2015, EN ISO13485:2016, EN1041:2008,

EN ISO14971:2012, EN ISO 15223-1:2016, EN ISO10993-5:2009, EN ISO 22523:2006

This declaration of conformity is issued under the sole responsibility of the manufacturer: OPPO Medical Inc., address at: Seattle City Center, 1420 Fifth Ave., Ste. 220080, Seattle, WA98101, USA

Oct. 12th, 2018

Date

President

EU Authorized Representative: MT Promedt Consulting GmbH

Altenhofstrasse 80 D-66386 St. Ingbert

Germany

MT Promedt Consulting GmbH (MTPC) has taken over the function and responsibilities of an European Authorized Representative of OPPO Medical Inc. in accordance to the requirements of the MDD 93/42/EEC.

The manufacturer has provided MTPC with the Technical Documentation.

The necessary notification of the above mentioned products has been performed. The products are registered under the Reg.-Nr.: DE/CA70/40838/0048119