

## EC – Declaration of Conformity

**Manufacturers Name:** Orfit Industries N.V.

**SRN (Single Registration Number):** BE-MF-000007872

**Manufacturers Address:** Vosveld 9a, 2110 Wijnegem, Belgium

**Basic UDI-DI:** 0542002870004MF

**Name of the Device(s):** Thermoplastic-moulding water bath - Suspan 1 & Suspan 2

**Product code(s):** 35097/230/1400, 35097/115/1400, 32502/230EU, 35097/RE

**Classification:** Class I, according the rules of Annex VIII

**Conformity assessment route:** Orfit Industries N.V. uses the following procedures for the CE-labelling of their products according the Regulation MDR 2017/745:

Class I: EC conformity declaration according to Annex IV.

**Applied norms:** ISO 13485:2016  
ISO 14971:2019  
ISO 15223-1:2021  
EN 55011 - 2011  
EN 55011 - 2018 (Class B)  
EN 60601-1-2  
EN 61000 - 3  
EN 61000 - 4  
EN 61010 -1

This declaration of conformity is issued under the sole responsibility of Orfit Industries N.V. We hereby declare that the medical device(s) specified above meet the provisions of the Regulation (EU) MDR 2017/745 for medical devices. This declaration is supported by the Quality System approval ISO 13485:2016 issued by Lloyd's Register EMEA – LRQA.

All supporting documentation is retained at the premises of the manufacturer.

Eddy Marivoet,  
Wijnegem, 10 August 2021



*Quality Assurance & Regulatory Affairs Manager*