

Declaration of Conformity

**Type of device:****Tissue Flootation Bath****Model:****TFB 35**

We hereby declare that the device named above has been designed to comply with all applicable essential requirements of the listed directives of the European Parliament:

EU Directive 98/79/EC
EU Directive 2014/30/EU

In Vitro Diagnostic Medical Devices
Electromagnetic compatibility (EMC)

according to:

EN 61010-2-101:2017-10
EN ISO 14971:2012
EN 61326-2-6:2013

Safety Requirements for IVD equipment
Risk Management
EMC for IVD equipment

Burgdorf, 02.01.2020



Dr. Michael Ott – Managing Director