CE

SLEE medical

Declaration of Conformity

We herewith declare, in exclusive responsibility, that the instruments

Tissue Processor MTP, MTM, MTM II
Paraffin Wax Embedding Centre MPS/P1, MPS/P2, MPS/P, MPS/W, MPS/C, MPS/CX
Rotary Microtome CUT4062, CUT5062, CUT6062
aquatec
Sledge Microtome cuttec S
Cryostat MTC, MEV, MEV plus, MNT
Staining Machine MSM, MAS
Linear Stainer cromatec
Cytocentrifuge CS I / II
Cover Slipper MCS I / II
printtec s & c, autoloader
Floating Bath / Slide Warmer: slidetec WATER, slidetec HEAT, slidetec WATER/HEAT, slidetec WATER s

are developed, designed and manufactured in compliance with the applicable directives

- **Council Directive 2014/35/EU, (Low Voltage) and**
- **Council Directive 2014/30/EU (Electromagnetic Compatibility)**
- **Directive 2011/65/EU (restriction of the use of certain hazardous substances in electrical and electronic equipment)**

including their amendments up to the date mentioned below.

The following harmonized standards were applied:

• **EN 61010-1:2010**

Safety requirements for electrical equipment for measurement, control and laboratory use -
Part 1: General requirements

• **EN 61326-1:2013**

Electrical equipment for measurement, control and laboratory use -
EMC requirements -
Part 1: General requirements

• **EN 61000-3-2:2014**

Electromagnetic compatibility (EMC)
Part 3-2: Limits - Limits for harmonic current emissions

• **EN 61000-3-3:2013**

Electromagnetic compatibility (EMC)
Part 3: Limits -
Section 3: Limitation of voltage fluctuations and flicker in low-voltage
supply systems for equipment with rated current ≤ 16 A

In addition, the following in-house standards were applied: **DIN EN ISO 9001:2015; EN ISO 13485:2016**

Mainz, 03.04.2019

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