CE

Leica MICROSYSTEMS

EC Declaration of Conformity

We herewith declare, in exclusive responsibility, that the instrument

Leica RM2125 RT – Rotary Microtome

was developed, designed and manufactured to conform with the

- Direktive 98/79/EC of the European Parliament and of the Council (in-vitro diagnostic medical devices)
- Directive 2006/42/EC of the European Parliament and of the Council on machinery

The following harmonized standards were applied:

- **DIN EN ISO 12100-1: 2003**
  Safety of machinery.
  Basic concepts, general principles for design.
  Part 1: Basic terminology, methodology

- **DIN EN ISO 12100-2: 2003**
  Safety of machinery.
  Basic concepts, general principles for design.
  Part 2: Technical principles and specifications.

In addition, the following in-house standards were applied:

- **DIN EN ISO 9001: 2000.**
  Quality management systems - Requirements

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