Advancing Cancer Diagnostics
Improving Lives

Leica
BIOSYSTEMS

EU DECLARATION OF CONFORMITY

We, **Leica Biosystems Nussloch GmbH**, **Heidelberger Str. 17-19**, **69226 Nussloch**, **Germany**

declare on our own responsibility, that the device

Leica HI1220

complies with

- Directive 2014/35/EU of the European Parliament and of the Council of 26 February 2014 on the harmonisation of the laws of the Member States relating to the making available on the market of electrical equipment designed for use within certain voltage limits (OJ L 96, 29.3.2014, p. 357–374)

  EN 61010-1:2010

- Directive 2014/30/EU of the European Parliament and of the Council of 26 February 2014 on the harmonisation of the laws of the Member States relating to electromagnetic compatibility (OJ L 96, 29.3.2014, p. 79–106)

  EN 61326-1:2013

- Directive 2011/65/EU of the European Parliament and of the Council of 8 June 2011 on the restriction of the use of certain hazardous substances in electrical and electronic equipment (OJ L 174, 1.7.2011, p. 88–110)

  EN 50581:2012

Quality Management System:	Certified according to EN ISO 13485:2016 and ISO 9001:2015
Manufacturing site:	Leica Microsystems Ltd. Shanghai, Building 1, 258 Jinzang Road, China (Shanghai) Pilot Free Trade, 201206 Shanghai, People's Republic of China

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Nussloch, 15.03.2019

ppa. Martin Laszak
Director Customer Service Core Histology

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Rev. A