

## **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 IP S

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

Nussloch, 15.03.2019

Wartin Laszak

Director Customer Service Core Histology

RA/QA Manager

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