

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 CM3600 XP

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

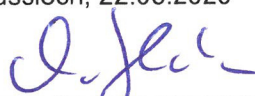
- Directive 2011/65/EU of the European Parliament and of the Council of 8 June 2011 on the restriction of the use of certain hazardous substances in electrical and electronic equipment (OJ L 174, 1.7.2011, p. 88–110)

EN 50581:2012

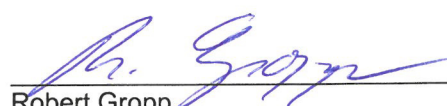
Quality Management System: Certified according to EN ISO 9001:2015

Manufacturing site: Leica Biosystems Nussloch GmbH, Heidelberger Str. 17-19,
69226 Nussloch, Germany

Nussloch, 22.06.2020



Andreas Hahn
Senior Director Core Histology



Robert Gropp
RA/QA Manager

Rev. B