

EU DECLARATION OF CONFORMITY

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

declare on our own responsibility, that the medical device

Leica ST5020

complies with

- the essential requirements of the Directive 98/79/EC of the European Parliament and of the Council of 27 October 1998 on in vitro diagnostic medical devices (OJ L 331, 7.12.1998, p. 1-37), Annex III

EN 61010-2-101:2017
EN ISO 14971:2012
EN 61326-2-6:2013

Device classification: IVD general/other

- 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, 07.05.2018


Martin Laszak
Director Customer Service Core Histology


Robert Gropp
RA/QA Manager