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 RM2245

complies with


  EN 61010-2-101:2002
  EN ISO 14971:2012
  EN 61326-2-6:2013

  Device classification: IVD general/other


  EN 50581:2012


Manufacturing site: Leica Microsystems Ltd, Shanghai, Building 1, 258 JinZang Road, Jinqiao Export&Processing Zone, Pudong, Shanghai 201206, PRC

Nussloch, 18.09.2017

Martin Laszak
Director Customer Service Core Histology & Site Leader Nussloch

Robert Gropp
RA/QA Manager