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 RM2255

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

Mátin Laszak
Director Customer Service Core Histology & Site Leader Nussloch

Robert Gropp
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