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

ASP6025 S

commits with


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

  Device classification: IVD general/other


  EN 50581:2012


Manufacturing sites: Leica Microsystems Ltd. Shanghai, Building 1, 258 Jinzang Road, China (Shanghai) Pilot Free Trade, 201206 Shanghai, People's Republic of China

Nussloch, 21.01.2020

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