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 ASP300 S

complies with

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

  Device classification: IVD general/other

  
  EN 50581:2012


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

Rev. A