EC 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 – Automated Vacuum Tissue Processor

meets 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 Annex III

This EC Declaration of Conformity is valid till 01.02.2013.

Nussloch, 21.02.2012

Arnd Kaldowski
President Biosystems Division