



Declaration of Conformity

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Conformity Assessment Procedure IVD Directive 98/79/EC, Annex III

Product(s):	Product Name	Cat. No.
	BD MAX™ UVE Sample Buffer Tubes	443420

We hereby declare that the above mentioned product(s) comply with the European In Vitro Diagnostic Directive and its relevant transposition into national laws of the member states into which we place the devices.

Date January 20, 2017

Name and Authority Bradford M Spring, VP Regulatory Affairs & Compliance
Privacy Officer
BD Diagnostic Systems and Preanalytical Systems

Signature