



Declaration of Conformity

Manufacturer:	Becton Dickinson and Company 7 Loveton Circle Sparks, Maryland 21152, USA Tel: +1.410.316.4000 Fax: +1.410.316.4499				
Authorized Representative:	Benex Limited Pottery Road, Dun Laoghaire Co. Dublin, Ireland Tel: +353.1.202.5222				
Conformity Assessment Procedure:	Directive 98/79/EC of the European Parliament and of the Council, Annex III of Directive 98/79/EC				
Product:	<table border="1"><tr><th>REF</th><th>Product Name</th></tr><tr><td>256045</td><td>BD Veritor™ System for Rapid Detection of Flu A+B</td></tr></table>	REF	Product Name	256045	BD Veritor™ System for Rapid Detection of Flu A+B
REF	Product Name				
256045	BD Veritor™ System for Rapid Detection of Flu A+B				
We hereby declare that the above-mentioned product complies with the European In Vitro Diagnostic Medical Devices Directive and its relevant transposition into national laws of the member states into which we place the devices. This declaration is issued under the sole responsibility of Becton, Dickinson and Company.					
Date:	07-Jan-2022				
Name and Authority:	Anne Zavertnik Vice President Regulatory Affairs, IDS				
Signature:					

RECORD REVISION HISTORY TABLE

Revision	Description of Changes
A	Initial Release in SAP