



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><tr><th>REF</th><th>Product Name</th></tr><tr><td>446252</td><td>BD Affirm™ VPIII Microbial Identification Test</td></tr><tr><td>446257</td><td>BD Affirm™ VPIII Microbial Identification Test</td></tr></table>	REF	Product Name	446252	BD Affirm™ VPIII Microbial Identification Test	446257	BD Affirm™ VPIII Microbial Identification Test
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446252	BD Affirm™ VPIII Microbial Identification Test						
446257	BD Affirm™ VPIII Microbial Identification Test						
We hereby declare that the above mentioned products comply 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:	02-Feb-2022						
Name and Authority:	Anne Zavertnik Vice President Regulatory Affairs, IDS						
Signature:							

RECORD REVISION HISTORY TABLE	
Revision	Description of Changes
A	Initial Release