



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>245122</td><td>BD BBL™ MGIT™ Mycobacteria Growth Indicator Tubes</td></tr></table>	REF	Product Name	245122	BD BBL™ MGIT™ Mycobacteria Growth Indicator Tubes
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245122	BD BBL™ MGIT™ Mycobacteria Growth Indicator Tubes				
We hereby declare that the above mentioned product 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: 20-Jan-2022					
Name and Authority:	Anne Zavertnik Vice President Regulatory Affairs, IDS				
Signature:					

Technical File Number: BALTER245122

RECORD REVISION HISTORY TABLE	
Revision	Description of Changes
A	Initial Release