



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												
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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"><thead><tr><th>REF</th><th>Product Name</th></tr></thead><tbody><tr><td>245123</td><td>BACTEC™ MGIT™ 960 SIRE Kit</td></tr><tr><td>245125</td><td>BD BACTEC™ MGIT™ 960 STR 4.0 Kit</td></tr><tr><td>245126</td><td>BD BACTEC™ MGIT™ 960 INH 0.4 Kit</td></tr><tr><td>245127</td><td>BD BACTEC™ MGIT™ 960 EMB 7.5 Kit</td></tr><tr><td>245157</td><td>BD BACTEC™ MGIT™ 960 IR Kit</td></tr></tbody></table>	REF	Product Name	245123	BACTEC™ MGIT™ 960 SIRE Kit	245125	BD BACTEC™ MGIT™ 960 STR 4.0 Kit	245126	BD BACTEC™ MGIT™ 960 INH 0.4 Kit	245127	BD BACTEC™ MGIT™ 960 EMB 7.5 Kit	245157	BD BACTEC™ MGIT™ 960 IR Kit
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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:	20-Jan-2022												
Name and Authority:	Anne Zavertnik Vice President Regulatory Affairs, IDS												
Signature:													

Technical File Number: BALTER245123

RECORD REVISION HISTORY TABLE	
Revision	Description of Changes
A	Initial Release