



Declaration of Conformity

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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>440928</td><td>BD ProbeTec™ Urine Preservative Transport Kit</td></tr><tr><td>440455</td><td>BD ProbeTec™ ET Sample Tubes and Caps</td></tr></tbody></table>	REF	Product Name	440928	BD ProbeTec™ Urine Preservative Transport Kit	440455	BD ProbeTec™ ET Sample Tubes and Caps
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We hereby declare that the above mentioned products 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. This declaration is issued under the sole responsibility of Becton, Dickinson and Company.							
Date: 22 DEC 2021							
Name and Authority:	Anne Zavertnik Vice President Regulatory Affairs, IDS						
Signature:							

RECORD REVISION HISTORY TABLE	
Revision	Description of Changes
A	Initial Release