



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 IV Notified Body: BSI Group The Netherlands, 2797						
Product:	<table><thead><tr><th>REF</th><th>Product Name</th></tr></thead><tbody><tr><td>441126</td><td>BD ProbeTec™ Chlamydia trachomatis (CT) Qx Amplified DNA Assay Reagent Pack</td></tr><tr><td>442959</td><td>BD ProbeTec™ Chlamydia trachomatis (CT) Qx Assay Gray Amp Reagent Pack</td></tr></tbody></table>	REF	Product Name	441126	BD ProbeTec™ Chlamydia trachomatis (CT) Qx Amplified DNA Assay Reagent Pack	442959	BD ProbeTec™ Chlamydia trachomatis (CT) Qx Assay Gray Amp Reagent Pack
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We hereby declare that the above mentioned products comply with the IVD Directive(98/79/EC) 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:	2021-Mar-9						
Name and Authority:	Kay Taylor Vice President Regulatory Affairs, Life Sciences						
Signature:							

Technical File Number: BDDSTF441126

RECORD REVISION HISTORY TABLE

Revision	Description of Changes
A	Initial SAP DoC release. Convert existing to new format.