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:

REF	Product Name
441125	Control Set for the BD ProbeTec™ Chlamydia trachomatis / Neisseria gonorrhoeae(CT/GC) Q^X Amplified DNA Assays
441925	Control Set for the BD ProbeTec™ Chlamydia trachomatis / Neisseria gonorrhoeae / Trichomonas vaginalis (CT/GC/TV) Q^X Amplified DNA Assays

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
WW Vice President Regulatory Affairs, IDS

Signature:

Technical File Number: BDDSTF441125

RECORD REVISION HISTORY TABLE	
Revision	**Description of Changes**
A	Initial Release