Declaration of Conformity

Manufacturer:	Becton, Dickinson and Company 7 Loveton Circle Sparks, MD 21152 USA
Authorized Representative:	Benex Limited Pottery Road, Dun Laoghaire Co. Dublin, Ireland Tel: +353.1.202.5222 Fax: +353.1.202.5388
Conformity assessment procedure:	Annex IV (Full Quality Assurance System) of the IVD Directive 98/79/EC, Notified Body: BSI 2797, Certificate Number: CE 614165
Product:	440450 - BD ProbeTec ET Chlamydia trachomatis (CT) Neisseria gonorrhoeae (GC) Amplification Control (AC) 440704 - BD ProbeTec ET Chlamydia trachomatis (CT) 440474 - BD ProbeTec ET Chlamydia trachomatis (CT) Amplification Control (AC) 440705 - BD ProbeTec ET Chlamydia trachomatis (CT) Neisseria gonorrhoeae (GC) 440451 - BD ProbeTec ET Control Set 440452 - BD ProbeTec ET Diluent (CT/GC) 440453 - BD ProbeTec ET Diluent (CT/GC)

We hereby declare that the above mentioned product(s) manufactured after 10/6/2019 complies 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.

Signed In Baltimore:	10/6/2019
Name and Authority:	Bradford M. Spring , VP, Regulatory Affairs
Signature:	

Technical File Number: BDDSTF440450