BD Declaration of Conformity

Manufacturer:	Becton Dickinson and Company 7 Loveton Circle Sparks, MD 21152, USA Tel: 410-316-4000 Fax: 410-316-4499
Authorized Representative:	Becton Dickinson France S.A.S. 11 rue Artistide Bergès 38800 Le Pont de Claix, France Tel: 04-76-68-3636 Fax: 04-76-68-3292
Conformity Assessment Procedure:	Council Directive 93/42/EEC as amended by 2007/47/EC, Annex VII, Section 5 for Class I products
Notified Body:	BSi (Notified Body number 2797) Say Building, John M. Keynesplein 9 1066 EP Amsterdam, Netherlands
Product:	220142 BD ProbeTec ET CT/GC Amplified DNA Assay Collection Kit for Endocervical Specimens 220143 BD ProbeTec ET CT/GC Amplified DNA Assay Collection Kit for Male Urethral Specimens 440476 BD ProbeTec ET CT/GC Amplified DNA Assay Endocervical Specimen Collection and Dry Transport Kit 440461 BD ProbeTec ET CT/GC Amplified DNA Assay Male Urethral Specimen Collection and Dry Transport Kit 441358 Male Urethral Specimen Collection Kit for the BD ProbeTec™ *Chlamydia trachomatis/Neisseria gonorrhoeae* (CT/GC) Q^x Amplified DNA Assays

We hereby declare that the above mentioned product(s) manufactured after 2009/10/21 complies with the European Medical Devices Directive and its relevant transposition into national laws of the member states into which we place the devices.

Signed in Baltimore:	2019/03/27
Name and Authority:	Bradford M. Spring Director, Regulatory Affairs
Signature: