Declaration of Conformity

Manufacturer:	Becton, Dickinson and Company 7 Loveton Circle Sparks, MD 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 Fax: +353.1.202.5388
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
443878	BD MAX™ MDR-TB

We hereby declare that the above mentioned product(s) manufactured after7/24/2019 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

Date:	January 26, 2021
Name and Authority:	Kay Taylor Vice President Regulatory Affairs, Life Sciences and IDS

Signature: Kay A Taylor

Technical File Number: <BDDSTF443878>

RECORD REVISION HISTORY TABLE	
Revision	**Description of Changes**
A	Initial SAP release. Transition to current format.