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
442020	BD BACTEC™ Peds Plus™/F Culture vials
442021	BD BACTEC™ Lytic/10 Anaerobic/F Culture Vials
442022	BD BACTEC™ Plus Anaerobic/F Culture Vials
442023	BD BACTEC™ Plus Aerobic/F Culture Vials
442024	BD BACTEC™ Standard Anaerobic/F Culture Vials (Plastic)
442027	BD BACTEC™ Standard/10 Aerobic/F Culture Vials (Plastic)
442017	BD BACTEC™ Mycosis IC/F Culture Vials (Plastic)

We hereby declare that the above mentioned products comply 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: 22 DEC 2021

Name and Authority:	Anne Zavertnik Vice President Regulatory Affairs, IDS

Signature: 

Technical File Number: BALTERBACTECPLASTIC

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