

## **Declaration of Conformity**

Becton Dickinson and Company

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Conformity
Assessment
Procedure:

**Product:** 

Directive 98/79/EC of the European Parliament and of the

Council. Annex III of Directive 98/79/EC

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:

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Status: Released EFFECTIVE
Revision: N/A Change #: N/A

Change #: N/A Classification: Confidential

RECORD REVISION HISTORY TABLE	
Revision	Description of Changes
Α	Initial Release

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