

BALTER232097-DOC(A)

Declaration of Conformity

Becton Dickinson and Company

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Conformity **Assessment** Procedure:

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

Council. Annex III of Directive 98/79/EC

REF Product Name

BD BBLTM Sensi-DiscTM Mupirocin 200 µg **Product:** 232097

We hereby declare that the above mentioned products comply with the European In Vitro Diagnostic Directive and their 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: 03-December-2021

Anne Zavertnik

Name and Authority: Vice President Regulatory Affairs, IDS

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RECORD REVISION HISTORY TABLE	
Revision	Description of Changes
A	Initial Release

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