

Manufacturer:

Declaration of Conformity

Becton Dickinson and Company

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Conformity Assessment Procedure:

Authorized

Representative:

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

Annex III of Directive 98/79/EC

Product:

REF	Product Name
445011	BD SARS-CoV-2/Flu for BD MAX™ System

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: November 13, 2020

Kay A. Taylor Name and Authority:

Vice President Regulatory Affairs, Life Sciences and IDS

Signature:

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RECORD REVISION HISTORY TABLE	
Revision	Description of Changes
01	Initial Release

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