



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 RoHS Directive 2011/65/EU of the European Parliament and of the Council as amended by Delegated Directive (EU) 2015/863, Annex II.									
Product:	<table border="1"> <thead> <tr> <th>REF</th> <th>Product Name</th> </tr> </thead> <tbody> <tr> <td>256066</td> <td>BD Veritor™ Plus System Analyzer</td> </tr> <tr> <td>256068</td> <td>BD Veritor™ InfoScan Module</td> </tr> <tr> <td>445010</td> <td>BD Veritor™ InfoWiFi Module</td> </tr> </tbody> </table>		REF	Product Name	256066	BD Veritor™ Plus System Analyzer	256068	BD Veritor™ InfoScan Module	445010	BD Veritor™ InfoWiFi Module
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<p>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.</p>										
Date:	December 23, 2021									
Name and Authority:	Anne Zavertnik Vice President Regulatory Affairs BD Integrated Diagnostic Solutions									
Signature:										

Technical File Number: BDDSTF256066

RECORD REVISION HISTORY TABLE	
Revision	Description of Changes
A	Initial Release
B	Software revision history change, change to GMDN code assigned to Veritor InfoWiFi module SKU, Modified section 5 to claim RoHS3 compliance, Provided justification for NA entries in the ERC checklist, migration to new template