



Declaration of Conformity

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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 -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>441007</td> <td>BD EpiCenter™ System Software</td> </tr> <tr> <td>440981</td> <td>BD EpiCenter™ Multi User Module</td> </tr> <tr> <td>440887</td> <td>BD EpiCenter™ Plus Software</td> </tr> </tbody> </table>	REF	Product Name	441007	BD EpiCenter™ System Software	440981	BD EpiCenter™ Multi User Module	440887	BD EpiCenter™ Plus Software
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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:	July 19, 2021								
Name and Authority:	Anne Zavertrnik Vice President Regulatory Affairs, IDS								
Signature:									

RECORD REVISION HISTORY TABLE	
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Revision	Description of Changes
A	Initial Release