



Declaration of Conformity

Manufacturer:	BD Kiestra B.V. Marconilaan 6 9207 JC Drachten The Netherlands Tel: +31.512. 510.710				
Authorized Representative:	N/A				
Conformity Assessment Procedure:	Directive 98/79/EC of the European Parliament and of the Council, Annex III of Directive 98/79/EC				
Product:	<table><tr><th>REF</th><th>Product Name</th></tr><tr><td>444910</td><td>BD Kiestra™ Methicillin-resistant Staphylococcus aureus (MRSA) Application Autorelease</td></tr></table>	REF	Product Name	444910	BD Kiestra™ Methicillin-resistant Staphylococcus aureus (MRSA) Application Autorelease
REF	Product Name				
444910	BD Kiestra™ Methicillin-resistant Staphylococcus aureus (MRSA) Application Autorelease				
We hereby declare that the above mentioned product comply with the 98/79/EC 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:	10 May 2021				
Name and Authority:	Karin Brands, Manager Regulatory Affairs				
Signature:					

RECORD REVISION HISTORY TABLE	
Revision	Description of Changes
A	New