



EU DECLARATION OF CONFORMITY (DoC)

Manufacturer:	Becton, Dickinson and Company 1 Becton Drive Franklin Lakes, New Jersey 07417 USA
Manufacturer SRN:	US-MF-000019182
Authorised Representative:	Becton Dickinson Ireland Ltd. Donore Road Co. Louth Drogheda, A92 YW26, Ireland
Authorised Representative SRN:	IE-AR-000007610
Product:	BD PosiFlush™ XS Syringes
Basic UDI-DI:	038290WKCQDZQWJK
Risk Class and Rule:	Class III, Annex VIII, Rule 14

Intended Purpose	<p>BD PosiFlush™ XS Syringes are intended to be used FOR FLUSHING ONLY of in-situ peripheral intravenous catheters (PIVCs), peripherally inserted central catheters (PICCs), central venous catheters (CVCs), and implanted venous access ports.</p> <p>BD PosiFlush™ XS Syringe is not intended for dry product reconstitution, for medication dilution, or where intravenous therapy with sodium chloride is indicated.</p> <p>Using aseptic technique, BD PosiFlush™ XS Syringe can be used on a sterile field.</p>
Notified Body:	<p>National Standards Authority of Ireland (NSAI)</p> <p>1, Swift square</p> <p>Northwood, Santry</p> <p>Dublin 9, Ireland</p> <p>Identification number : 0050</p>
<p>We, as the manufacturer of the device(s) take sole responsibility for and hereby declare that the above mentioned product(s) meet(s) the provisions of the following Directives/ Regulation(s):</p> <ul style="list-style-type: none"> Regulation (EU) 2017/745 of the European Parliament and of the Council on Medical Devices 	

Conformity Assessment Route:

<input checked="" type="checkbox"/> ANNEX IX Chapter I and III – Quality Management System	<p>EC CERTIFICATE No.: 745.008</p> <p>Certificate Expiration Date: 20 December 2027</p>
<input checked="" type="checkbox"/> ANNEX IX Chapter II – Technical Documentation	<p>EC CERTIFICATE No.: 745.008</p> <p>Certificate Expiration Date: 20 December 2027</p>
<input type="checkbox"/> ANNEX X Type Examination	<p>EC CERTIFICATE No.:</p> <p>Certificate Expiration Date:</p>
<input type="checkbox"/> ANNEX XI Part A Production Quality Assurance	<p>EC CERTIFICATE No.:</p> <p>Certificate Expiration Date:</p>
<input type="checkbox"/> ANNEX XI Part B Product Verification	<p>EC CERTIFICATE No.:</p> <p>Certificate Expiration Date:</p>
<input type="checkbox"/> ANNEX II & III Technical Documentation	N/A

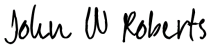

Common Specifications (CS): Common Specifications have not been issued for this product.

Number: <Version/Year>	Title:	Full or Partial Application: <Justification>
N/A	N/A	N/A

**Devices Covered by this DoC:**

SKU#	Device Name	Device Class
306570	BD PosiFlush™ XS syringe CE 3mL	III
306571	BD PosiFlush™ XS syringe CE 5mL	III
306572	BD PosiFlush™ XS syringe CE 10mL	III
306580	BD PosiFlush™ XS syringe EMA 3mL	III
306581	BD PosiFlush™ XS syringe EMA 5mL	III
306582	BD PosiFlush™ XS syringe EMA 10mL	III

Authorised Signatory:

Name & Title:	John W. Roberts Regulatory Affairs Director Medication Delivery Solutions
On behalf of:	Becton, Dickinson and Company
Place of Issue:	
Date of Issue:	
Signature:	<div>DocuSigned by:  8B97BB78BFD485...</div> <div>DS </div>

DECLARATION OF CONFORMITY Revision History:

Version:	Date:	Detailed Change Description:	Prepared by:
A	22 December 2022	New document created to meet MDR (EU) 2017/745 compliance.	Perrine Clert-Girard