## **BD Preanalytical Systems - Regulatory Affairs Procedure**



Document Number: VTF0003-02 Revision Level: 04

TITLE: Declaration of Conformity for BD Vacutainer® Luer Adapter

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## **EC DECLARATION OF CONFORMITY**

Legal Manufacturer:	Becton Dickinson and Company (BD) 1 Becton Drive Franklin Lakes, NJ 07417 USA
Authorised Representative:	Becton Dickinson and Company Belliver Industrial Estate Belliver Way Roborough Plymouth PL6 7BP UK
Manufacturing Site(s):	Becton Dickinson and Company 1575 Airport Road Sumter, SC 29153
Products:	367300 BD Vacutainer® Luer Adapter
Classification:	EU Class I Sterile per Annex IX, Rule 1 of the Medical Device Directive (93/42/EEC).  Canada Class I per Schedule 1, Part 1, Rule 7, sub-rule 1 of the Canadian Medical Device Regulations (CMDR), SOR//98-282 which states that all other non-invasive devices are classified as Class I in which none of the indents apply.
Conformity Assessment Route:	Annex II, Medical Device Directive 93/42/EEC
GMDN:	GMDN Code: 60579 GMDN Term: Blood collection luer adapter

We herewith declare that the above mentioned products meet the provisions of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices. All supporting documentation is retained at the premises of the manufacturer.

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Standards:	EN 556-1 – 2001
	EN 1041 - 2013
	EN-ISO-15223-1:2016
	EN ISO 11135 - 2014
	EN ISO 11607-1 – 2010
	EN ISO 11737-1 – 2006
	EN ISO 13485 – 2016
	EN ISO 14971 – 2012
	EN ISO 14155 - 2011
Notified Body:	National Standards Association of Ireland (NSAI)
100	1 Swift Square
	Northwood
	Santry, Dublin 9, Ireland
	Phone: 353 (01) 807-3800
	Fax: 353 (01) 807-3838
CE Certificate Number:	252-548
Date of issuance of original CE certificate:	19 May 1997

Date: 27 April 2018

Vernon Brown

**Director Regulatory Affairs** 

**BD Preanalytical Systems** 

Becton, Dickinson and Company (BD)

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REVISION HISTORY			
Current Version Prepared By: Pamela Sanecki			
REV.	Revision Description	Releasing ECO (if applicable)	
01	Initial Release	N/A	
02	Review and released – no updates necessary.	N/A	
03	Changed GMDN code to 60579 as the old number was obsoleted. In the Standards section, deleted EN-980 and updated the revision date on EN ISO 15223-1-2016.	N/A	
04	Updated Standards revision dates to comply with V08-510-01 rev. 04.	N/A	