



# Quality System Approval Certificate

Medical Devices Directive 93/42/EEC

*The National Standards Authority of Ireland as a duly designated  
Notified Body, (identification number 0050), for the purposes of the European Communities  
(Medical Devices) Regulations (S.I. No. 252 of 1994)*

*APPROVES THE QUALITY SYSTEM APPLIED BY*

## Becton, Dickinson and Company Limited

**Donore Road  
Drogheda  
Co Louth  
Ireland**

*to the Product Family*

### Syringes (BD Plastipak™ and BD™ Perfusion)

**GMDN Code: 47017**

*on the basis of examination under the requirements of Directive 93/42/EEC on Medical Devices Annex V.  
The use of the NSAI Notified Body identification number 0050 in conjunction with CE Marking of  
Conformance for this product family is hereby authorized.*

<b>Registration Number:</b>	<b>252.156</b>
<b>Original Approval:</b>	<b>13 September 1995</b>
<b>Last Amended on:</b>	<b>12 September 2016</b>
<b>Remains valid until:</b>	<b>12 September 2019</b>

**Signed:**

Approved by:  
Kevin D. Mullaney  
Chief Executive Officer - NSAI Inc.

Approved by:  
Susan Murphy  
European Medical Device Operations Manager

**This certificate remains valid on condition that the Approved Quality System is maintained in an adequate and efficacious manner .**  
Details of the current product range and operational locations included within the scope of this approval can be obtained from NSAI

**National Standards Authority of Ireland, 1 Swift Square, Northwood, Santry, Dublin 9, Ireland.**