



# 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

**1 Becton Drive  
Franklin Lakes  
NJ 07417  
USA**

*to the Product Family*

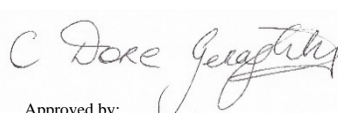
**Pharmacy Products - Medicine administration kit, oral; Medication  
transfer needle, filtering; Medicine preparation needle/cannula; Syringe  
tip cap**


**GMDN Code: 16266, 16627, 63614, 64357, 64514**

*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 authorised.*

<b>Registration Number:</b>	<b>252.308</b>
<b>Original Approval:</b>	<b>02 April 1998</b>
<b>Last Amended on:</b>	<b>24 May 2021</b>
<b>Remains valid until:</b>	<b>03 February 2024</b>

**Signed:**

  
Approved by:  
Dr. Caroline Dore Geraghty  
Director, Medical Devices

  
Approved by:  
Dr. Elaine Darcy  
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.**