



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

**Hypodermic Syringes, insulin and general use (BD Micro-Fine™ +,
BD Micro-Fine™ Plus, Micro-Fine™ IV, Ultra-Fine™ and Ultra-
Fine™ II Insulin Syringes and Plastipak™ Allergy Syringes)**

GMDN Code: 38501, 35904

*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.*

Registration Number:	252.140
Original Approval:	07 April 1995
Last Amended on:	15 April 2020
Remains valid until:	25 May 2024

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.