

K924070 B-D PEN NEEDLEMar 18, 1993
218 days to decisionK924070 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k924070/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Aug 12, 1992
Decision date	Mar 18, 1993
Days to decision	218 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Bd Becton Dickinson Vacutainer Systems Preanalytic
Location	Washington, DC, US
Contact	PETER ZURLO
510(k) history	632 submissions · 625 cleared · 1976-2001

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k924070/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 21, 2026