FDA 510(k) Premarket Notification

K924070 B-D PEN NEEDLE

CLEARED

Mar 18, 1993
218 days to decision

K924070 · Product code: **FMI** · General Hospital
Source: 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