FDA 510(k) Premarket Notification

K924439 BECTON DICKINSON INTERLINK VIAL ACCESS CANNULA

CLEARED

Feb 12, 1993
164 days to decision

K924439 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k924439/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Sep 1, 1992
Decision date	Feb 12, 1993
Days to decision	164 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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