FDA 510(k) Premarket Notification

K912330 B-D(TM) MICROFINE(TM) IV PEN INJECTOR CAT. #328204

CLEARED

Oct 11, 1991
140 days to decision

K912330 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k912330/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	May 24, 1991
Decision date	Oct 11, 1991
Days to decision	140 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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