FDA 510(k) Premarket Notification

K901756 INTERJECTOR BLOOD COLLECTION DEVICE

CLEARED

Aug 29, 1990
134 days to decision

K901756 · Product code: **JKA** · Chemistry
Source: https://www.510kdatabase.net/k901756/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Vials, Systems, Serum Separators, Blood Collection (JKA)
Date received	Apr 17, 1990
Decision date	Aug 29, 1990
Days to decision	134 days
Third-party review	No

APPLICANT

Company	**Bowman/Vector, Inc.**
Location	Fort Lauderdale, FL, US
Contact	BLAIR, PA
510(k) history	1 submissions · 1 cleared · 1990-1990