

K901756 INTERJECTOR BLOOD COLLECTION DEVICEAug 29, 1990
134 days to decisionK901756 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k901756/>; 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 June 30, 2026