

**K964654 BECTON DICKINSON BLUNT PLASTIC CANNULA**Apr 2, 1997  
133 days to decisionK964654 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k964654/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Nov 20, 1996
Decision date	Apr 2, 1997
Days to decision	133 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Bd Becton Dickinson Vacutainer Systems Preanalytic</b>
Location	Washington, DC, US
Contact	GREGORY W MORGAN
510(k) history	632 submissions · 625 cleared · 1976-2001

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k964654/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 21, 2026