

**K924439 BECTON DICKINSON INTERLINK VIAL ACCESS  
CANNULA**Feb 12, 1993  
164 days to decisionK924439 · Product code: **FMI** · General Hospital  
Source: <https://www.510kdatabase.net/k924439/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k924439/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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