

**K912330 B-D(TM) MICROFINE(TM) IV PEN INJECTOR CAT.  
#328204**Oct 11, 1991  
140 days to decisionK912330 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k912330/>**SUBMISSION DETAILS**

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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**

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Company	<b>Bd Becton Dickinson Vacutainer Systems Preanalytic</b>
Location	Washington, DC, US
Contact	J ARNSBERGER
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/k912330/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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