

**K972264 FIRST MIDCATH CATHETER**Sep 15, 1997  
90 days to decisionK972264 · Product code: **FOZ** · General HospitalSource: <https://www.510kdatabase.net/k972264/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	Jun 17, 1997
Decision date	Sep 15, 1997
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Becton Dickinson Infusion Therapy Systems, Inc.</b>
Location	Sandy, UT, US
Contact	C.J. WELLE
510(k) history	36 submissions · 36 cleared · 1997-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k972264/> 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 26, 2026