

**K031600 CONDUCTION CATHETER**Aug 13, 2003  
84 days to decisionK031600 · Product code: **FRN** · General Hospital  
Source: <https://www.510kdatabase.net/k031600/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion (FRN)
Date received	May 21, 2003
Decision date	Aug 13, 2003
Days to decision	84 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Micor, Inc.</b>
Location	Pittsburgh, PA, US
Contact	SUSAN FORGRAVE
510(k) history	8 submissions · 8 cleared · 1986-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k031600/> 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 25, 2026