

**K890751 ACCI-GUARD(TM) I.V. CATHETER PLACEMENT UNIT**Apr 21, 1989  
67 days to decisionK890751 · Product code: **FOZ** · General HospitalSource: <https://www.510kdatabase.net/k890751/>**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	Feb 13, 1989
Decision date	Apr 21, 1989
Days to decision	67 days
Third-party review	No

**APPLICANT**

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Company	<b>Medical Safety Products, Inc.</b>
Location	Denver, CO, US
Contact	E SAGSTETTER
510(k) history	6 submissions · 6 cleared · 1988-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k890751/>, 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 June 30, 2026