

**K833092 DESERET ANGIO-GUIDE LARGE VEIN CATH.**Nov 21, 1983  
70 days to decisionK833092 · Product code: **FOZ** · General Hospital  
Source: <https://www.510kdatabase.net/k833092/>**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	Sep 12, 1983
Decision date	Nov 21, 1983
Days to decision	70 days
Third-party review	No

**APPLICANT**

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Company	<b>Warner-Lambert Co.</b>
Location	Mchenry, IL, US
510(k) history	50 submissions · 50 cleared · 1979-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k833092/> 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 14, 2026