

**K821577 DESERET ANGIO GUIDE**Jun 16, 1982  
19 days to decisionK821577 · Product code: **FOZ** · General HospitalSource: <https://www.510kdatabase.net/k821577/>**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	May 28, 1982
Decision date	Jun 16, 1982
Days to decision	19 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/k821577/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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