

K821372 DESERET INTRACATH INTRAVENOUS CATHETERJun 2, 1982
23 days to decisionK821372 · Product code: **FOZ** · General Hospital
Source: <https://www.510kdatabase.net/k821372/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	May 10, 1982
Decision date	Jun 2, 1982
Days to decision	23 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k821372/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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