

**K910799 LANDMARK VENOUS ACCESS DEVICE**Mar 21, 1991  
24 days to decisionK910799 · Product code: **FOZ** · General HospitalSource: <https://www.510kdatabase.net/k910799/>**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 25, 1991
Decision date	Mar 21, 1991
Days to decision	24 days
Third-party review	No

**APPLICANT**

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Company	<b>Menlo Care, Inc.</b>
Location	Menlo Park, CA, US
Contact	JOCK M WALKER
510(k) history	31 submissions · 25 cleared · 1986-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k910799/>; 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 July 2, 2026