

**K890435 VITAACCESS CATHETER INTRODUCER KIT**Mar 29, 1989  
61 days to decisionK890435 · Product code: **FOZ** · General Hospital  
Source: <https://www.510kdatabase.net/k890435/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - SD
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	Jan 27, 1989
Decision date	Mar 29, 1989
Days to decision	61 days
Third-party review	No

**APPLICANT**

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Company	<b>Vitaphore Corp.</b>
Location	San Francisco, CA, US
Contact	JESSICA AYRES
510(k) history	13 submissions · 12 cleared · 1986-1992

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k890435/>; 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 22, 2026