

**K000632 0.9% SODIUM CHLORIDE VASCULAT ACCESS  
DEVICE**Oct 19, 2000  
238 days to decisionK000632 · Product code: **FOZ** · General Hospital  
Source: <https://www.510kdatabase.net/k000632/>**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 24, 2000
Decision date	Oct 19, 2000
Days to decision	238 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Avitro, LLC</b>
Location	Woodstock, IL, US
Contact	JOHN BRDA
510(k) history	2 submissions · 2 cleared · 2000-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k000632/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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