

**K914528 SAFEDWEL(R) CENTRAL VENOUS CATHETER**Oct 13, 1992  
370 days to decisionK914528 · Product code: **FOZ** · General HospitalSource: <https://www.510kdatabase.net/k914528/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - KD
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	Oct 9, 1991
Decision date	Oct 13, 1992
Days to decision	370 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Becton Dickinson Vascular Access, Inc.</b>
Location	Sandy, UT, US
Contact	WELLE
510(k) history	25 submissions · 22 cleared · 1992-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k914528/> 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 5, 2026