

**K860290 VIAPIC CENTRAL LINE CATHETER**Mar 18, 1986  
49 days to decisionK860290 · Product code: **FOZ** · General HospitalSource: <https://www.510kdatabase.net/k860290/>**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	Jan 28, 1986
Decision date	Mar 18, 1986
Days to decision	49 days
Third-party review	No

**APPLICANT**

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Company	<b>Parke-Davis Co.</b>
Location	Mchenry, IL, US
Contact	E. F. CLEARY
510(k) history	47 submissions · 47 cleared · 1976-1986

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k860290/>, 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 27, 2026