

**K900320 CRI CYNOSAR CATHETER**Sep 14, 1990  
234 days to decisionK900320 · Product code: **DRA** · CardiovascularSource: <https://www.510kdatabase.net/k900320/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Steerable (DRA)
Date received	Jan 23, 1990
Decision date	Sep 14, 1990
Days to decision	234 days
Third-party review	No

**APPLICANT**

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Company	<b>Catheter Research C/O Burditt, Bowles &amp; Radzius</b>
Location	Chicago, IL, US
Contact	BRIAN GRIGSBY
510(k) history	8 submissions · 8 cleared · 1987-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k900320/>, 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 6, 2026