

**K896690 MODIFIED ANGIOGRAPHIC CATHETER**Jan 24, 1990  
57 days to decisionK896690 · Product code: **DQO** · CardiovascularSource: <https://www.510kdatabase.net/k896690/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Nov 28, 1989
Decision date	Jan 24, 1990
Days to decision	57 days
Third-party review	No

**APPLICANT**

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Company	<b>Novoste Corp.</b>
Location	Miami, FL, US
Contact	CHARLES LARSEN
510(k) history	5 submissions · 5 cleared · 1988-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k896690/>, 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 June 28, 2026