

**K770430 CATHETER, CURVED LEFT HEART VENT**Mar 16, 1977  
9 days to decisionK770430 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k770430/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Mar 7, 1977
Decision date	Mar 16, 1977
Days to decision	9 days
Third-party review	No

**APPLICANT**

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Company	<b>3M Health Care, Sarns</b>
Location	Mchenry, IL, US
510(k) history	76 submissions · 76 cleared · 1976-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k770430/> 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 1, 2026