

**K792152 CARDIOPLEGIA SOLUTION ADMIN. SET**Nov 13, 1979  
22 days to decisionK792152 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k792152/>**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	Oct 22, 1979
Decision date	Nov 13, 1979
Days to decision	22 days
Third-party review	No

**APPLICANT**

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Company	<b>Cardio Systems, Inc.</b>
Location	Mchenry, IL, US
510(k) history	9 submissions · 9 cleared · 1979-2013

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k792152/> 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