

**K896636 RMI ANTEPLEGIA CANNULA**Feb 15, 1990  
85 days to decisionK896636 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k896636/>**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	Nov 22, 1989
Decision date	Feb 15, 1990
Days to decision	85 days
Third-party review	No

**APPLICANT**

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Company	<b>Research Medical, Inc.</b>
Location	Mchenry, IL, US
Contact	MICHAEL N KELLY
510(k) history	35 submissions · 35 cleared · 1984-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k896636/>; 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