

**K870109 RMI VENOUS DRAINAGE CANNULA**Mar 16, 1987  
63 days to decisionK870109 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k870109/>**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	Jan 12, 1987
Decision date	Mar 16, 1987
Days to decision	63 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/k870109/>; 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 7, 2026