

**K861322 RMI POROUS VENT PLUG**May 30, 1986  
52 days to decisionK861322 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k861322/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Apr 8, 1986
Decision date	May 30, 1986
Days to decision	52 days
Third-party review	No

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k861322/> 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