

**K960098 RMI INTERNAL MAMMARY ARTERY CANNULA**Mar 26, 1996  
75 days to decisionK960098 · Product code: **DQR** · Cardiovascular  
Source: <https://www.510kdatabase.net/k960098/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Catheter (DQR)
Date received	Jan 11, 1996
Decision date	Mar 26, 1996
Days to decision	75 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k960098/> 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