

**K951088 QUEST RETROGRADE CARDIPLÉGIA CANNULA-  
RCCS**Jun 14, 1995  
97 days to decisionK951088 · Product code: **DWF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k951088/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Mar 9, 1995
Decision date	Jun 14, 1995
Days to decision	97 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Quest Medical, Inc.</b>
Location	Walker, MI, US
Contact	DREW JOHNSON
510(k) history	39 submissions · 39 cleared · 1980-2024

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k951088/> Data sourced from FDA 510(k) public records (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. - Generated June 28, 2026