

**K042685 METRICATH SYSTEM**Nov 18, 2004  
50 days to decisionK042685 · Product code: **DQO** · Cardiovascular  
Source: <https://www.510kdatabase.net/k042685/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Sep 29, 2004
Decision date	Nov 18, 2004
Days to decision	50 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Angiometrx, Inc.</b>
Location	Seattle, WA, US
Contact	TIM VERSPAGEN
510(k) history	3 submissions · 3 cleared · 2003-2005

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k042685/> 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 2, 2026