

**K990173 PATHFINDER II ANGIOGRAPHIC CATHETER**Dec 21, 1999  
336 days to decisionK990173 · Product code: **DQO** · CardiovascularSource: <https://www.510kdatabase.net/k990173/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Jan 19, 1999
Decision date	Dec 21, 1999
Days to decision	336 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Maxxim Medical</b>
Location	Arlington, TX, US
Contact	EDDIE MONROE
510(k) history	26 submissions · 25 cleared · 1994-2002

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k990173/> 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 June 29, 2026