

**K880887 NEOCATH**May 24, 1988  
83 days to decisionK880887 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k880887/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Mar 2, 1988
Decision date	May 24, 1988
Days to decision	83 days
Third-party review	No

**APPLICANT**

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Company	<b>Taris Corp.</b>
Location	Fernandina Beach, FL, US
Contact	WILLIAM J SCHRODER
510(k) history	3 submissions · 3 cleared · 1988-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k880887/>; 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 7, 2026