

**K872788 MODIFIED CAROTID SHUNT W/SPRING REINFORCED  
LUMEN**Sep 25, 1987  
73 days to decisionK872788 · Product code: **DWF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k872788/>**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	Jul 14, 1987
Decision date	Sep 25, 1987
Days to decision	73 days
Third-party review	No

**APPLICANT**

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Company	<b>Uresil Corp.</b>
Location	Mchenry, IL, US
Contact	MICHAEL JARON
510(k) history	45 submissions · 44 cleared · 1981-2001

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k872788/>; 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 22, 2026