

**K881455 EMBOLECTOMY CATHETER(MODIFIED)**Jun 15, 1988  
70 days to decisionK881455 · Product code: **DXE** · CardiovascularSource: <https://www.510kdatabase.net/k881455/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Embolectomy (DXE)
Date received	Apr 6, 1988
Decision date	Jun 15, 1988
Days to decision	70 days
Third-party review	No

**APPLICANT**

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Company	<b>Cathlab Corp.</b>
Location	Tustin, CA, US
Contact	SATTAR SAUDAGAR
510(k) history	8 submissions · 8 cleared · 1988-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k881455/>; 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