

**K880571 CATH-TECH PORT IMPLANTABLE VASCULAR ACCESS SYSTEM**Mar 4, 1988  
23 days to decisionK880571 · Product code: LJT · General Hospital  
Source: <https://www.510kdatabase.net/k880571/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intravascular (LJT)
Date received	Feb 10, 1988
Decision date	Mar 4, 1988
Days to decision	23 days
Third-party review	No

**APPLICANT**

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Company	<b>Catheter Technology Corp.</b>
Location	Walker, MI, US
Contact	JACK SPEER
510(k) history	13 submissions · 13 cleared · 1983-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k880571/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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