

**K882230 A PORT IMPLANT VASCULAR ACCESS SYSTEM -  
DETACHED**Jul 29, 1988  
63 days to decisionK882230 · Product code: **LJT** · General Hospital  
Source: <https://www.510kdatabase.net/k882230/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intravascular (LJT)
Date received	May 27, 1988
Decision date	Jul 29, 1988
Days to decision	63 days
Third-party review	No

**APPLICANT**

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Company	<b>Therex Corp.</b>
Location	Walpole, MA, US
Contact	BRADLEY J ENEGREN
510(k) history	26 submissions · 23 cleared · 1988-1995

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k882230/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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