

**K901419 DUAL PORT**Sep 5, 1990  
163 days to decisionK901419 · Product code: **LJT** · General Hospital  
Source: <https://www.510kdatabase.net/k901419/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intravascular (LJT)
Date received	Mar 26, 1990
Decision date	Sep 5, 1990
Days to decision	163 days
Third-party review	No

**APPLICANT**

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Company	<b>Infusaid Corp.</b>
Location	Walker, MI, US
Contact	NANCY C HALL
510(k) history	8 submissions · 8 cleared · 1983-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k901419/>; 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 6, 2026