

**K841679 VASCULAR-ACCESS-PORT WITHDRAWAL-BLOOD**Jun 28, 1984  
66 days to decisionK841679 · Product code: **LJT** · General Hospital  
Source: <https://www.510kdatabase.net/k841679/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intravascular (LJT)
Date received	Apr 23, 1984
Decision date	Jun 28, 1984
Days to decision	66 days
Third-party review	No

**APPLICANT**

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Company	<b>Norfolk Medical Products, Inc.</b>
Location	Walker, MI, US
510(k) history	20 submissions · 20 cleared · 1983-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k841679/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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