

K830000 VASCULAR-ACCESS-PORTMar 1, 1983
57 days to decisionK830000 · Product code: **LJT** · General Hospital
Source: <https://www.510kdatabase.net/k830000/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intravascular (LJT)
Date received	Jan 3, 1983
Decision date	Mar 1, 1983
Days to decision	57 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k830000/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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