

**K942784 NORFOLK OMEGA PORT-LP SUBCUTANEOUSLY
IMPLANTED**May 2, 1995
322 days to decisionK942784 · Product code: **LJT** · General Hospital
Source: <https://www.510kdatabase.net/k942784/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intravascular (LJT)
Date received	Jun 14, 1994
Decision date	May 2, 1995
Days to decision	322 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k942784/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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