FDA 510(k) Premarket Notification

K942784 NORFOLK OMEGA PORT-LP SUBCUTANEOUSLY IMPLANTED

CLEARED

May 2, 1995
322 days to decision

K942784 · 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