FDA 510(k) Premarket Notification

K921201 VITAL-PORT M.R.I. DUAL LUMEN VASCULAR ACCESS SYST.

CLEARED

Jul 21, 1992
131 days to decision

K921201 · Product code: **LJT** · General Hospital
Source: https://www.510kdatabase.net/k921201/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intravascular (LJT)
Date received	Mar 12, 1992
Decision date	Jul 21, 1992
Days to decision	131 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Med Institute, Inc.**
Location	West Lafayette, IN, US
Contact	NEAL E FEARNOT
510(k) history	26 submissions · 23 cleared · 1990-2000