

K921201 VITAL-PORT M.R.I. DUAL LUMEN VASCULAR ACCESS SYST.Jul 21, 1992
131 days to decisionK921201 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k921201/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 20, 2026