

K030636 C-PORTJun 5, 2003
97 days to decisionK030636 · Product code: **LJT** · General Hospital
Source: <https://www.510kdatabase.net/k030636/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intravascular (LJT)
Date received	Feb 28, 2003
Decision date	Jun 5, 2003
Days to decision	97 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Clinisurg Vascular Products, LLC
Location	Dallas, TX, US
Contact	DONALD F HULTS
510(k) history	1 submissions · 1 cleared · 2003-2003

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k030636/> 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 July 2, 2026