

K934504 HMP DUAL TITANIUM PORTFeb 25, 1994
163 days to decisionK934504 · Product code: **LJT** · General Hospital
Source: <https://www.510kdatabase.net/k934504/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent - K
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intravascular (LJT)
Date received	Sep 15, 1993
Decision date	Feb 25, 1994
Days to decision	163 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Horizon Medical Products, Inc.
Location	Atlanta, GA, US
Contact	RODDY J H. CLARK
510(k) history	16 submissions · 11 cleared · 1994-2003

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k934504/> 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 June 30, 2026