

**K934640 HMP DUAL POLYSULFONE PORT**Feb 25, 1994  
150 days to decisionK934640 · Product code: **LJT** · General Hospital  
Source: <https://www.510kdatabase.net/k934640/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - K
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intravascular (LJT)
Date received	Sep 28, 1993
Decision date	Feb 25, 1994
Days to decision	150 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k934640/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 30, 2026