

**K001874 PERIPHERALLY INSERTED CENTRAL CATHETER**Sep 25, 2000  
97 days to decisionK001874 · Product code: **LJS** · General Hospital  
Source: <https://www.510kdatabase.net/k001874/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Long-term Greater Than 30 Days (LJS)
Date received	Jun 20, 2000
Decision date	Sep 25, 2000
Days to decision	97 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Utah Medical Products, Inc.</b>
Location	Mchenry, IL, US
Contact	KEVIN L CORNWELL
510(k) history	38 submissions · 38 cleared · 1979-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k001874/> 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 May 23, 2026