

**K031718 LIFEVALVE CENTRAL VENOUS CATHTER**Jul 2, 2003  
29 days to decisionK031718 · Product code: **LJS** · General Hospital  
Source: <https://www.510kdatabase.net/k031718/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Intravascular, Therapeutic, Long-term Greater Than 30 Days (LJS)
Date received	Jun 3, 2003
Decision date	Jul 2, 2003
Days to decision	29 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	SCOTT MOELLER
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/k031718/> 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