

**K921755 8 FR SINGLE LUMEN HIGH PERF. GROSHONG CENT.  
VENOUS**Oct 23, 1992  
193 days to decisionK921755 · Product code: **LJS** · General Hospital  
Source: <https://www.510kdatabase.net/k921755/>**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	Apr 13, 1992
Decision date	Oct 23, 1992
Days to decision	193 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Davol, Inc.</b>
Location	Mchenry, IL, US
Contact	JACK SPEER
510(k) history	50 submissions · 47 cleared · 1977-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k921755/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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