

K083763 ZEUS CT PICC 4 FR AND 5 FR SINGLE AND DUAL LUMENMar 16, 2009
88 days to decisionK083763 · Product code: **LJS** · General Hospital
Source: <https://www.510kdatabase.net/k083763/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Long-term Greater Than 30 Days (LJS)
Date received	Dec 18, 2008
Decision date	Mar 16, 2009
Days to decision	88 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	R4 Vascular, Inc.
Location	St. Paul, MN, US
Contact	LAURIE LEWANDOWSKI
510(k) history	7 submissions · 5 cleared · 2007-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k083763/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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