

**K990713 SPECTRANETICS LEAD LOCKING DEVICE (LLD) #1,  
#2, #3 LISTER**

Oct 22, 1999  
232 days to decision

K990713 · Product code: **DRB** · Cardiovascular  
Source: <https://www.510kdatabase.net/k990713/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stylet, Catheter (DRB)
Date received	Mar 4, 1999
Decision date	Oct 22, 1999
Days to decision	232 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Spectranetics Corp.</b>
Location	Colorado Springs, CO, US
Contact	MICHAEL J QUINN
510(k) history	24 submissions · 24 cleared · 1999-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k990713/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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