

**K032411 JS VASCULAR GUIDE WIRE VISE/TORQUE DEVICE**Oct 7, 2003  
63 days to decisionK032411 · Product code: **DQX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k032411/>**SUBMISSION DETAILS**

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|                       |                                    |
|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Traditional                        |
| Device classification | Wire, Guide, Catheter (DQX)        |
| Date received         | Aug 5, 2003                        |
| Decision date         | Oct 7, 2003                        |
| Days to decision      | 63 days                            |
| Third-party review    | No                                 |
| Summary / Statement   | Summary                            |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>Js Vascular, Inc.</b>              |
| Location       | Orinda, CA, US                        |
| Contact        | MICHAEL A DANIEL                      |
| 510(k) history | 1 submissions · 1 cleared · 2003-2003 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k032411/> 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 July 3, 2026