

**K860430 THE NAVIGATOR STEERABLE GUIDE WIRE**Jul 18, 1986  
164 days to decisionK860430 · Product code: **DQX** · CardiovascularSource: <https://www.510kdatabase.net/k860430/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Feb 4, 1986
Decision date	Jul 18, 1986
Days to decision	164 days
Third-party review	No

**APPLICANT**

---

Company	<b>Madox Surgimed, Inc.</b>
Location	Mchenry, IL, US
Contact	JANE AOYAGI
510(k) history	24 submissions · 24 cleared · 1984-1995

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k860430/>, 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 1, 2026