

**K860429 MEADOX SURGIMED GUIDE CATHETER**Apr 23, 1986  
78 days to decisionK860429 · Product code: **DQO** · CardiovascularSource: <https://www.510kdatabase.net/k860429/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Feb 4, 1986
Decision date	Apr 23, 1986
Days to decision	78 days
Third-party review	No

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k860429/> 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