

**K854912 ANGIOMED GUIDED WIRES**Apr 16, 1986  
128 days to decisionK854912 · Product code: **DQX** · CardiovascularSource: <https://www.510kdatabase.net/k854912/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Dec 9, 1985
Decision date	Apr 16, 1986
Days to decision	128 days
Third-party review	No

**APPLICANT**

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Company	<b>Angiomed U.S., Inc.</b>
Location	Anaheim, CA, US
Contact	RICHARD P MOHR
510(k) history	24 submissions · 24 cleared · 1986-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k854912/>; 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 June 29, 2026