

**K991898 ULTRA-SELECT GUIDEWIRE, HYTEK GUIDEWIRE**Jun 16, 1999  
12 days to decisionK991898 · Product code: **DQX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k991898/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Wire, Guide, Catheter (DQX)
Date received	Jun 4, 1999
Decision date	Jun 16, 1999
Days to decision	12 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Microvena Corp.</b>
Location	Findley, MN, US
Contact	ANGELA MALLERY
510(k) history	18 submissions · 18 cleared · 1990-1999

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k991898/> 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 4, 2026