

**K822597 GUIDE WIRE**Sep 21, 1982  
25 days to decisionK822597 · Product code: **DQX** · CardiovascularSource: <https://www.510kdatabase.net/k822597/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Aug 27, 1982
Decision date	Sep 21, 1982
Days to decision	25 days
Third-party review	No

**APPLICANT**

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Company	<b>Vertex Medical Corp.</b>
Location	Mchenry, IL, US
510(k) history	21 submissions · 21 cleared · 1982-1984

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k822597/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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