

**K830420 POLYETHYLENE MONITORING CONNECTORS**Feb 28, 1983  
20 days to decisionK830420 · Product code: **DQO** · CardiovascularSource: <https://www.510kdatabase.net/k830420/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Feb 8, 1983
Decision date	Feb 28, 1983
Days to decision	20 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/k830420/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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