

**K832064 VENOUS CATHETER**Aug 8, 1983  
42 days to decisionK832064 · Product code: **DQY** · Cardiovascular  
Source: <https://www.510kdatabase.net/k832064/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Jun 27, 1983
Decision date	Aug 8, 1983
Days to decision	42 days
Third-party review	No

**APPLICANT**

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Company	<b>Cardio Search, Inc.</b>
Location	Walker, MI, US
510(k) history	4 submissions · 4 cleared · 1983-1985

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

Device record: <https://www.510kdatabase.net/k832064/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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