

**K822243 GUIDING CATHETER**Aug 24, 1982  
28 days to decisionK822243 · Product code: **DQO** · CardiovascularSource: <https://www.510kdatabase.net/k822243/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Jul 27, 1982
Decision date	Aug 24, 1982
Days to decision	28 days
Third-party review	No

**APPLICANT**

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Company	<b>Interventional Medical, Inc.</b>
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1982-1988

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

Device record: <https://www.510kdatabase.net/k822243/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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