

**K813577 PRESSURE INDICATOR-REGULATOR**Jan 29, 1982  
37 days to decisionK813577 · Product code: **DQO** · CardiovascularSource: <https://www.510kdatabase.net/k813577/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Dec 23, 1981
Decision date	Jan 29, 1982
Days to decision	37 days
Third-party review	No

**APPLICANT**

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Company	<b>Advanced Catheter Systems</b>
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1982-1982

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k813577/> 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 9, 2026