

**K831636 PERCUTANEOUS INTRODUCER KIT**Aug 24, 1983  
93 days to decisionK831636 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k831636/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	May 23, 1983
Decision date	Aug 24, 1983
Days to decision	93 days
Third-party review	No

**APPLICANT**

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Company	<b>Accu-Line</b>
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1983-1983

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

Device record: <https://www.510kdatabase.net/k831636/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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