

**K881088 ADDITION OF A TIP GUARD TO THE  
ATHEROCATH(TM)**May 31, 1988  
77 days to decisionK881088 · Product code: LIT · Cardiovascular  
Source: <https://www.510kdatabase.net/k881088/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Angioplasty, Peripheral, Transluminal (LIT)
Date received	Mar 15, 1988
Decision date	May 31, 1988
Days to decision	77 days
Third-party review	No

**APPLICANT**

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Company	<b>Devices For Vascular Intervention, Inc.</b>
Location	Redwood City, CA, US
Contact	DIANE RUPPERT
510(k) history	14 submissions · 14 cleared · 1987-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k881088/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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