

**K893658 PERCUTANEOUS INTRODUCER**Aug 3, 1989  
80 days to decisionK893658 · Product code: **DYB** · CardiovascularSource: <https://www.510kdatabase.net/k893658/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	May 15, 1989
Decision date	Aug 3, 1989
Days to decision	80 days
Third-party review	No

**APPLICANT**

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Company	<b>Medamicus, Inc.</b>
Location	Minneapolis, MN, US
Contact	DENNIS MADISON
510(k) history	20 submissions · 20 cleared · 1989-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k893658/>; 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 June 28, 2026