

**K870907 INTRODUCER, PERCUTANEOUS**Apr 9, 1987  
35 days to decisionK870907 · Product code: **DYB** · CardiovascularSource: <https://www.510kdatabase.net/k870907/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Mar 5, 1987
Decision date	Apr 9, 1987
Days to decision	35 days
Third-party review	No

**APPLICANT**

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Company	<b>Med Fusion Systems, Inc.</b>
Location	Mchenry, IL, US
Contact	RUFFIN BOOTH
510(k) history	22 submissions · 22 cleared · 1977-1987

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k870907/>; 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 1, 2026