

**K871153 CATHETER INTRODUCER**Jun 24, 1987  
92 days to decisionK871153 · Product code: **DYB** · CardiovascularSource: <https://www.510kdatabase.net/k871153/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Mar 24, 1987
Decision date	Jun 24, 1987
Days to decision	92 days
Third-party review	No

**APPLICANT**

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Company	<b>Angeion Corp.</b>
Location	Plymouth, MN, US
Contact	LAWRENCE M KANE
510(k) history	7 submissions · 7 cleared · 1987-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k871153/>; 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 2, 2026