

**K903906 PERCUTANEOUS CATHETER INTRODUCER KIT**Sep 26, 1990  
34 days to decisionK903906 · Product code: **DYB** · CardiovascularSource: <https://www.510kdatabase.net/k903906/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Aug 23, 1990
Decision date	Sep 26, 1990
Days to decision	34 days
Third-party review	No

**APPLICANT**

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Company	<b>Intec Medical, Inc.</b>
Location	Blue Springs, MO, US
Contact	KENNETH A SPECTOR
510(k) history	17 submissions · 17 cleared · 1980-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k903906/>; 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 May 27, 2026