

**K072130 PTFE PEEL AWAY MICROINTRODUCER**Oct 31, 2007  
90 days to decisionK072130 · Product code: **DYB** · CardiovascularSource: <https://www.510kdatabase.net/k072130/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Aug 2, 2007
Decision date	Oct 31, 2007
Days to decision	90 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Intromed</b>
Location	Salt Lake City, UT, US
Contact	PHIL TRIOLO
510(k) history	1 submissions · 1 cleared · 2007-2007

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k072130/> 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 26, 2026