

**K051601 PINNACLE DESTINATION PERIPHERAL GUIDING SHEATH**Aug 4, 2005  
49 days to decisionK051601 · Product code: **DYB** · Cardiovascular  
Source: <https://www.510kdatabase.net/k051601/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Jun 16, 2005
Decision date	Aug 4, 2005
Days to decision	49 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Terumo Medical Corp.</b>
Location	Elkton, MD, US
Contact	MARK UNTERREINER
510(k) history	143 submissions · 143 cleared · 1980-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k051601/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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