

**K152173 Glidesheath**Dec 1, 2015  
119 days to decisionK152173 · Product code: **DYB** · CardiovascularSource: <https://www.510kdatabase.net/k152173/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Aug 4, 2015
Decision date	Dec 1, 2015
Days to decision	119 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Terumo Medical Corporation</b>
Location	Elkton, MD, US
Contact	ERIN DOYLE
510(k) history	14 submissions · 14 cleared · 2011-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k152173/> 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 26, 2026