

**K142183 GLIDESHEATH SLENDER**Nov 21, 2014  
105 days to decisionK142183 · Product code: **DYB** · Cardiovascular  
Source: <https://www.510kdatabase.net/k142183/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Introducer, Catheter (DYB)
Date received	Aug 8, 2014
Decision date	Nov 21, 2014
Days to decision	105 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/k142183/> 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