

**K122980 GLIDESHEATH SLENDER MODEL RM\* ES6J10HQS,  
GLIDESHEATH SLENDER MODEL RM\* ES6F16HQ,  
GLIDESHEATH SLENDER MODEL RM \* RS6J10PQ**Dec 11, 2012  
76 days to decisionK122980 · Product code: **DYB** · Cardiovascular  
Source: <https://www.510kdatabase.net/k122980/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Sep 26, 2012
Decision date	Dec 11, 2012
Days to decision	76 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Terumo Medical Corporation</b>
Location	Elkton, MD, US
Contact	DANIEL R PLONSKI
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/k122980/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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