

**K150542 Terumo Pump Tubing**May 1, 2015  
59 days to decisionK150542 · Product code: **DWE** · Cardiovascular  
Source: <https://www.510kdatabase.net/k150542/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Tubing, Pump, Cardiopulmonary Bypass (DWE)
Date received	Mar 3, 2015
Decision date	May 1, 2015
Days to decision	59 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Terumo Cardiovascular Systems Corporation</b>
Location	Elkton, MD, US
Contact	JOSHUA EWING
510(k) history	29 submissions · 29 cleared · 2002-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k150542/> 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 5, 2026