

**K072313 AMPLATZER TORQVUE DELIVERY SYSTEM**Nov 2, 2007  
77 days to decisionK072313 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k072313/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Aug 17, 2007
Decision date	Nov 2, 2007
Days to decision	77 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Aga Medical Corp.</b>
Location	Plymouth, MN, US
Contact	PATRICIA A LAFORTE
510(k) history	14 submissions · 14 cleared · 2000-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k072313/> 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 14, 2026