

**K051917 DIVER C.E. CATHETER**Aug 8, 2005  
24 days to decisionK051917 · Product code: **QEZ** · CardiovascularSource: <https://www.510kdatabase.net/k051917/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Aspiration Thrombectomy Catheter (QEZ)
Date received	Jul 15, 2005
Decision date	Aug 8, 2005
Days to decision	24 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Invatec Innovative Technologies, S.R.L.</b>
Location	Plymouth, MN, US
Contact	STEPHANIE K ISGRIGG ROBINSON
510(k) history	7 submissions · 7 cleared · 2005-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k051917/> 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 23, 2026