

**K140438 MICRO GUIDE CATHETER ELITE**Nov 14, 2014  
266 days to decisionK140438 · Product code: **PDU** · Cardiovascular  
Source: <https://www.510kdatabase.net/k140438/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter For Crossing Total Occlusions (PDU)
Date received	Feb 21, 2014
Decision date	Nov 14, 2014
Days to decision	266 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Cordis Corp., A Johnson &amp; Johnson Co.</b>
Location	Warren, NJ, US
Contact	Michelle Ragozzine Rodgers
510(k) history	5 submissions · 5 cleared · 2006-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k140438/> 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 28, 2026