

**K160848 Wingman 18 Crossing Catheter**Jul 14, 2016  
108 days to decisionK160848 · Product code: **DQY** · Cardiovascular  
Source: <https://www.510kdatabase.net/k160848/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Mar 28, 2016
Decision date	Jul 14, 2016
Days to decision	108 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Reflow Medical, Inc.</b>
Location	San Clemente, CA, US
Contact	Rebecca K Pine
510(k) history	9 submissions · 8 cleared · 2016-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k160848/> 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 26, 2026