

**K160121 CorPath GRX System**Oct 27, 2016  
282 days to decisionK160121 · Product code: **DXX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k160121/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Catheter Control, Steerable (DXX)
Date received	Jan 19, 2016
Decision date	Oct 27, 2016
Days to decision	282 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Corindus, Inc.</b>
Location	Baltimore, MD, US
Contact	Michael Douglas
510(k) history	9 submissions · 9 cleared · 2012-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k160121/> 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