

**K202275 CorPath GRX System**Dec 11, 2020  
122 days to decisionK202275 · Product code: **DXX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k202275/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Catheter Control, Steerable (DXX)
Date received	Aug 11, 2020
Decision date	Dec 11, 2020
Days to decision	122 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Corindus, Inc.</b>
Location	Baltimore, MD, US
Contact	Dana Hartlein
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/k202275/> 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