

**K141175 RENOVOCATH RC120**Oct 24, 2014  
171 days to decisionK141175 · Product code: **MJN** · Cardiovascular  
Source: <https://www.510kdatabase.net/k141175/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular Occluding, Temporary (MJN)
Date received	May 6, 2014
Decision date	Oct 24, 2014
Days to decision	171 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Renovorx, Inc.</b>
Location	Sunnyvale, CA, US
Contact	RON S WARREN
510(k) history	4 submissions · 4 cleared · 2014-2021

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k141175/> 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 25, 2026