

**K171290 Vela RX PTA Balloon Dilatation Catheter**Jul 19, 2017  
78 days to decisionK171290 · Product code: LIT · Cardiovascular  
Source: <https://www.510kdatabase.net/k171290/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Angioplasty, Peripheral, Transluminal (LIT)
Date received	May 2, 2017
Decision date	Jul 19, 2017
Days to decision	78 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Curatia Medical</b>
Location	Santa Clara, CA, US
Contact	Jessica Chiu
510(k) history	1 submissions · 1 cleared · 2017-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k171290/> 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 June 1, 2026