

**K133624 COMPASS CT, COMPASS CT PORT**Apr 18, 2014  
143 days to decisionK133624 · Product code: **DRS** · Cardiovascular  
Source: <https://www.510kdatabase.net/k133624/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	Nov 26, 2013
Decision date	Apr 18, 2014
Days to decision	143 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Mirador Biomedical</b>
Location	Seattle, WA, US
Contact	JUSTIN HULVERSHORN
510(k) history	1 submissions · 1 cleared · 2014-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k133624/> 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 7, 2026