

**K192802 primeMidline Catheters**Oct 28, 2019  
28 days to decisionK192802 · Product code: **PND** · General Hospital  
Source: <https://www.510kdatabase.net/k192802/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Midline Catheter (PND)
Date received	Sep 30, 2019
Decision date	Oct 28, 2019
Days to decision	28 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Pfm Medical, Inc.</b>
Location	Oceanside, CA, US
Contact	Mina Jiang
510(k) history	20 submissions · 17 cleared · 2004-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k192802/> 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 2, 2026