

**K890828 PERIFLUX PF3**Jun 20, 1989  
119 days to decisionK890828 · Product code: **DPW** · CardiovascularSource: <https://www.510kdatabase.net/k890828/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Flowmeter, Blood, Cardiovascular (DPW)
Date received	Feb 21, 1989
Decision date	Jun 20, 1989
Days to decision	119 days
Third-party review	No

**APPLICANT**

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Company	<b>Perimed, Inc.</b>
Location	Piscataway, NJ, US
Contact	OYSTEIN LIND
510(k) history	6 submissions · 6 cleared · 1989-1999

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k890828/>; 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