

**K844631 PERIFLUX PF-2**Feb 20, 1985  
85 days to decisionK844631 · Product code: **DPW** · CardiovascularSource: <https://www.510kdatabase.net/k844631/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Flowmeter, Blood, Cardiovascular (DPW)
Date received	Nov 27, 1984
Decision date	Feb 20, 1985
Days to decision	85 days
Third-party review	No

**APPLICANT**

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Company	<b>Medex, Inc.</b>
Location	Mchenry, IL, US
Contact	MARK T SANDRINI
510(k) history	48 submissions · 46 cleared · 1977-2005

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k844631/>; 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 29, 2026