

**K051231 VERIFYNOW-P2Y12 ASSAY**Aug 5, 2005  
84 days to decisionK051231 · Product code: **JOZ** · Hematology  
Source: <https://www.510kdatabase.net/k051231/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Automated Platelet Aggregation (JOZ)
Date received	May 13, 2005
Decision date	Aug 5, 2005
Days to decision	84 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Accumetrics, Inc.</b>
Location	San Diego, CA, US
Contact	BARBARA STEVENS
510(k) history	7 submissions · 7 cleared · 1999-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k051231/> 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 4, 2026