

**K980839 AVOCET PT**Sep 30, 1998  
210 days to decisionK980839 · Product code: **JPA** · Hematology  
Source: <https://www.510kdatabase.net/k980839/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Multipurpose For In Vitro Coagulation Studies (JPA)
Date received	Mar 4, 1998
Decision date	Sep 30, 1998
Days to decision	210 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Avocet Medical, Inc.</b>
Location	Campbell, CA, US
Contact	Judith Blunt
510(k) history	4 submissions · 4 cleared · 1998-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k980839/> 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 28, 2026