

**K083896 DESTINY MAX COAGULATION ANALYZER**Jul 2, 2009  
185 days to decisionK083896 · Product code: **JPA** · Hematology  
Source: <https://www.510kdatabase.net/k083896/>**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	Dec 29, 2008
Decision date	Jul 2, 2009
Days to decision	185 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Trinity Biotech</b>
Location	Jamestown, NY, US
Contact	Bonnie B DeJoy
510(k) history	5 submissions · 5 cleared · 2009-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k083896/> 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 31, 2026