

**K883635 KD 0.1 M SODIUM CITRATE SOLUTION**Oct 31, 1988  
67 days to decisionK883635 · Product code: **JPA** · Hematology  
Source: <https://www.510kdatabase.net/k883635/>**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	Aug 25, 1988
Decision date	Oct 31, 1988
Days to decision	67 days
Third-party review	No

**APPLICANT**

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Company	<b>King Diagnostics, Inc.</b>
Location	Walker, MI, US
Contact	FRANCES LOH
510(k) history	41 submissions · 41 cleared · 1981-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k883635/>; 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