

**K971236 GREINER VACUETTE BLOOD COLLECTION TUBE**May 12, 1997  
40 days to decisionK971236 · Product code: **JKA** · Toxicology  
Source: <https://www.510kdatabase.net/k971236/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Vials, Systems, Serum Separators, Blood Collection (JKA)
Date received	Apr 2, 1997
Decision date	May 12, 1997
Days to decision	40 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Greiner America, Inc.</b>
Location	Wilmington, DE, US
Contact	ED MAIER
510(k) history	7 submissions · 7 cleared · 1996-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k971236/> 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