

**K002777 VACUETTE WITH PPACK**Oct 5, 2000  
29 days to decisionK002777 · Product code: **JKA** · Chemistry  
Source: <https://www.510kdatabase.net/k002777/>**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	Sep 6, 2000
Decision date	Oct 5, 2000
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Greiner Vacuette North America, Inc.</b>
Location	Ellicott City, MD, US
Contact	Judi Smith
510(k) history	5 submissions · 5 cleared · 2000-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k002777/> 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