

**K014104 VACUETTE EDTA K2 TUBES**Feb 1, 2002  
50 days to decisionK014104 · Product code: **PJE** · Chemistry  
Source: <https://www.510kdatabase.net/k014104/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Blood/plasma Collection Device For Dna Testing (PJE)
Date received	Dec 13, 2001
Decision date	Feb 1, 2002
Days to decision	50 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/k014104/> 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