

**K030394 FLUID SAMPLE RETRIEVAL SYSTEM**Oct 27, 2003  
264 days to decisionK030394 · Product code: **JKA** · Chemistry  
Source: <https://www.510kdatabase.net/k030394/>**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	Feb 5, 2003
Decision date	Oct 27, 2003
Days to decision	264 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Biomed Personal Metabolic and Nutritional Testing</b>
Location	Memphis, TN, US
Contact	SUSAN B FENTRESS
510(k) history	2 submissions · 2 cleared · 2003-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k030394/> 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