

**K030076 PROFESSIONAL BLOOD SAMPLE RETRIEVAL SYSTEM**Jul 7, 2003  
180 days to decisionK030076 · Product code: **JKA** · Chemistry  
Source: <https://www.510kdatabase.net/k030076/>**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	Jan 8, 2003
Decision date	Jul 7, 2003
Days to decision	180 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/k030076/>; 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