

**K926123 DUAL CHAMBER MIXING BAG**Sep 3, 1993  
273 days to decisionK926123 · Product code: **KPE** · General Hospital  
Source: <https://www.510kdatabase.net/k926123/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Container, I.v. (KPE)
Date received	Dec 4, 1992
Decision date	Sep 3, 1993
Days to decision	273 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Secure Medical, Inc.</b>
Location	Mundelein, IL, US
Contact	TED D LYJAK
510(k) history	1 submissions · 1 cleared · 1993-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k926123/> 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 July 7, 2026