

**K930225 K-LOK CATHETER SECUREMENT DEVICE**Apr 21, 1993  
96 days to decisionK930225 · Product code: **KMK** · General Hospital  
Source: <https://www.510kdatabase.net/k930225/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Intravascular Catheter Securement (KMK)
Date received	Jan 15, 1993
Decision date	Apr 21, 1993
Days to decision	96 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>K-Lok, Inc.</b>
Location	Boca Raton, FL, US
Contact	GLENDAL KALT
510(k) history	2 submissions · 2 cleared · 1993-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k930225/> 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