

**K935252 K-LOK CATHETER SECUREMENT DEVICE  
MODIFICATION**Feb 18, 1994  
134 days to decisionK935252 · Product code: **KMK** · General Hospital  
Source: <https://www.510kdatabase.net/k935252/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Intravascular Catheter Securement (KMK)
Date received	Oct 7, 1993
Decision date	Feb 18, 1994
Days to decision	134 days
Third-party review	No
Summary / Statement	Statement

**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/k935252/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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