

**K843882 TRENDS LANCET**Nov 5, 1984  
34 days to decisionK843882 · Product code: **KMK** · General Hospital  
Source: <https://www.510kdatabase.net/k843882/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Intravascular Catheter Securement (KMK)
Date received	Oct 2, 1984
Decision date	Nov 5, 1984
Days to decision	34 days
Third-party review	No

**APPLICANT**

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Company	<b>Orange Medical Instruments, Inc.</b>
Location	Mchenry, IL, US
Contact	HELENE EHRLICH
510(k) history	12 submissions · 12 cleared · 1982-1986

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k843882/>; 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 May 27, 2026