

**K892440 KENDALL CURITY EXTENSION TUBES**Jul 24, 1989  
105 days to decisionK892440 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k892440/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Apr 10, 1989
Decision date	Jul 24, 1989
Days to decision	105 days
Third-party review	No

**APPLICANT**

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Company	<b>The Kendal Co.</b>
Location	Mchenry, IL, US
Contact	WILLIAM D CURTIN
510(k) history	63 submissions · 60 cleared · 1980-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k892440/>, 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 22, 2026