

**K854862 CURITY URETERAL STENT**Feb 10, 1986  
67 days to decisionK854862 · Product code: **FAD** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k854862/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stent, Ureteral (FAD)
Date received	Dec 5, 1985
Decision date	Feb 10, 1986
Days to decision	67 days
Third-party review	No

**APPLICANT**

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Company	<b>The Kendal Co.</b>
Location	Mchenry, IL, US
Contact	CHRISTINE MORGIDA
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/k854862/>; 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