

**K830884 SURGITEKS PERCUTANEOUS ANTEGRADE ALL -**May 9, 1983  
49 days to decisionK830884 · Product code: **FAD** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k830884/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stent, Ureteral (FAD)
Date received	Mar 21, 1983
Decision date	May 9, 1983
Days to decision	49 days
Third-party review	No

**APPLICANT**

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Company	<b>Medical Engineering Corp.</b>
Location	Mchenry, IL, US
510(k) history	28 submissions · 28 cleared · 1977-1993

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k830884/> 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