

**K813488 CATHETER REGULAR**Dec 31, 1981  
30 days to decisionK813488 · Product code: **FJS** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k813488/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Peritoneal, Long-term Indwelling (FJS)
Date received	Dec 1, 1981
Decision date	Dec 31, 1981
Days to decision	30 days
Third-party review	No

**APPLICANT**

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Company	<b>Accurate Surgical Instruments Co.</b>
Location	Walker, MI, US
Contact	ACCURATE SURGICAL INSTRUMENTS CO
510(k) history	9 submissions · 9 cleared · 1981-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k813488/>; 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 July 1, 2026