

K852389 URETERAL STENTSep 5, 1985
92 days to decisionK852389 · Product code: **FAD** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k852389/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stent, Ureteral (FAD)
Date received	Jun 5, 1985
Decision date	Sep 5, 1985
Days to decision	92 days
Third-party review	No

APPLICANT

Company	American Medical Systems, Inc.
Location	Mchenry, IL, US
Contact	MARK E MCINTYRE
510(k) history	72 submissions · 72 cleared · 1979-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k852389/>; Data sourced from FDA 510(k) public records (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. - Generated May 14, 2026