

K840059 SILITEK URINARY DIVERSION STENTFeb 4, 1984
26 days to decisionK840059 · Product code: **FAD** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k840059/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stent, Ureteral (FAD)
Date received	Jan 9, 1984
Decision date	Feb 4, 1984
Days to decision	26 days
Third-party review	No

APPLICANT

Company	Surgitek
Location	Mchenry, IL, US
510(k) history	29 submissions · 28 cleared · 1983-1995

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Device record: <https://www.510kdatabase.net/k840059/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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