

K833782 SURGITEK ALL SILICONE INFECTION DOUBJApr 2, 1984
157 days to decisionK833782 · Product code: **FAD** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k833782/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stent, Ureteral (FAD)
Date received	Oct 28, 1983
Decision date	Apr 2, 1984
Days to decision	157 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/k833782/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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