

K800109 SURGITEK URINARY DIVERSION STENTFeb 26, 1980
41 days to decisionK800109 · Product code: **FAD** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k800109/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stent, Ureteral (FAD)
Date received	Jan 16, 1980
Decision date	Feb 26, 1980
Days to decision	41 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k800109/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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