

**K855176 TRANSURETHAL PROSTATOME DIAG TREATM
PROPULSION SYS**May 13, 1986
138 days to decisionK855176 · Product code: **FBK** · Radiology
Source: <https://www.510kdatabase.net/k855176/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Endoscopic Injection Needle, Gastroenterology-urology (FBK)
Date received	Dec 26, 1985
Decision date	May 13, 1986
Days to decision	138 days
Third-party review	No

APPLICANT

Company	Urologic Surgical Devices Corp.
Location	Chicago, IL, US
Contact	HAROLD E COLLINS
510(k) history	1 submissions · 1 cleared · 1986-1986

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k855176/> 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 21, 2026