

**K890629 ESOPHAGEAL OVERTUBE**May 31, 1989  
110 days to decisionK890629 · Product code: **FED** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k890629/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Endoscopic Access Overtube, Gastroenterology-urology (FED)
Date received	Feb 10, 1989
Decision date	May 31, 1989
Days to decision	110 days
Third-party review	No

**APPLICANT**

---

Company	<b>Gastrotech, Inc.</b>
Location	Independence, OH, US
Contact	PAT GILPIN
510(k) history	5 submissions · 4 cleared · 1986-1989

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k890629/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated July 2, 2026