

K970892 FIEGERT ENDOTECH CYSTOSCOPEMay 5, 1997
55 days to decisionK970892 · Product code: **FAJ** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k970892/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cystoscope And Accessories, Flexible/rigid (FAJ)
Date received	Mar 11, 1997
Decision date	May 5, 1997
Days to decision	55 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Fiegert, Inc.
Location	Hollywood, FL, US
Contact	ERNESTO HERNANDEZ
510(k) history	1 submissions · 1 cleared · 1997-1997

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k970892/> 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 July 1, 2026