FDA 510(k) Premarket Notification

K970892 FIEGERT ENDOTECH CYSTOSCOPE

CLEARED

May 5, 1997
55 days to decision

K970892 · Product code: **FAJ** · Gastroenterology & Urology
Source: 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