

K945421 CHAKOFF ENDOSCOPYDec 23, 1994
49 days to decisionK945421 · Product code: **FDG** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k945421/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Retractor, Fiberoptic (FDG)
Date received	Nov 4, 1994
Decision date	Dec 23, 1994
Days to decision	49 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Stephen Chakoff, Inc.
Location	Miami, FL, US
Contact	FRANK GOLDFARB
510(k) history	16 submissions · 16 cleared · 1993-1995

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k945421/> 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 June 29, 2026