FDA 510(k) Premarket Notification

K945421 CHAKOFF ENDOSCOPY

CLEARED

Dec 23, 1994
49 days to decision

K945421 · Product code: **FDG** · Gastroenterology & Urology
Source: 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