FDA 510(k) Premarket Notification

K962212 KARL STORZ FRIMBERGER VARIOGUIDE

CLEARED

Dec 13, 1996
186 days to decision

K962212 · Product code: **FDT** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k962212/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Duodenoscope And Accessories, Flexible/rigid (FDT)
Date received	Jun 10, 1996
Decision date	Dec 13, 1996
Days to decision	186 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**KARL STORZ Endoscopy-America, Inc.**
Location	Mchenry, IL, US
Contact	MARIKA ANDERSON
510(k) history	361 submissions · 361 cleared · 1980-2024