

K962212 KARL STORZ FRIMBERGER VARIOGUIDEDec 13, 1996
186 days to decisionK962212 · Product code: **FDT** · Gastroenterology & UrologySource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k962212/> 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 May 15, 2026