

K962969 MEDICAL DEVICE TECHNOLOGIES, INC.ULTRADec 6, 1996
128 days to decisionK962969 · Product code: **KNW** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k962969/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy (KNW)
Date received	Jul 31, 1996
Decision date	Dec 6, 1996
Days to decision	128 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Medical Device Technologies, Inc.
Location	Gainesville, FL, US
Contact	KARL SWARTZ
510(k) history	46 submissions · 46 cleared · 1992-2010

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k962969/> 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 20, 2026