

K990839 TRU-CORE I REUSABLE BIOPSY INSTRUMENTMay 20, 1999
66 days to decisionK990839 · Product code: **KNW** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k990839/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy (KNW)
Date received	Mar 15, 1999
Decision date	May 20, 1999
Days to decision	66 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/k990839/> 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