FDA 510(k) Premarket Notification

K962969 MEDICAL DEVICE TECHNOLOGIES, INC.ULTRA

CLEARED

Dec 6, 1996
128 days to decision

K962969 · Product code: **KNW** · Gastroenterology & Urology
Source: 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	Gainsville, FL, US
Contact	KARL SWARTZ
510(k) history	46 submissions · 46 cleared · 1992-2010