FDA 510(k) Premarket Notification

K990839 TRU-CORE I REUSABLE BIOPSY INSTRUMENT

CLEARED

May 20, 1999
66 days to decision

K990839 · Product code: **KNW** · Gastroenterology & Urology
Source: 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	Gainsville, FL, US
Contact	KARL SWARTZ
510(k) history	46 submissions · 46 cleared · 1992-2010