

**K944837 MEDICAL DEVICE TECHNOLOGIES MANUAL BIOPSY  
NEEDLE**Oct 14, 1994  
14 days to decisionK944837 · Product code: **KNW** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k944837/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy (KNW)
Date received	Sep 30, 1994
Decision date	Oct 14, 1994
Days to decision	14 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k944837/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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