

**K921418 ULTRA-CORE BIOPSY NEEDLES**Jun 19, 1992  
87 days to decisionK921418 · Product code: **DWO** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k921418/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Biopsy, Cardiovascular (DWO)
Date received	Mar 24, 1992
Decision date	Jun 19, 1992
Days to decision	87 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	PAUL BAKER
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/k921418/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 20, 2026