

**K911448 MULTIPLE BIOPSY DEVICE-MBX**Apr 16, 1991  
32 days to decisionK911448 · Product code: **FCK** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k911448/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy, Suction (FCK)
Date received	Mar 15, 1991
Decision date	Apr 16, 1991
Days to decision	32 days
Third-party review	No

**APPLICANT**

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Company	<b>Triton Technology, Inc.</b>
Location	San Diego, CA, US
Contact	WILLIAM Z KOLOZSI
510(k) history	4 submissions · 4 cleared · 1988-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k911448/>; 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 June 29, 2026