

**K034021 VACUUM ASSISTED CORE BIOPSY DEVICE**Sep 8, 2004  
259 days to decisionK034021 · Product code: **KNW** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k034021/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy (KNW)
Date received	Dec 24, 2003
Decision date	Sep 8, 2004
Days to decision	259 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Suros Surgical Systems, Inc.</b>
Location	Indianapolis, IN, US
Contact	JOSEPH L MARK
510(k) history	3 submissions · 3 cleared · 2004-2007

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k034021/> 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