

**K924665 QUICKANCHOR**Oct 27, 1992  
47 days to decisionK924665 · Product code: **JDR** · Orthopedic  
Source: <https://www.510kdatabase.net/k924665/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Staple, Fixation, Bone (JDR)
Date received	Sep 10, 1992
Decision date	Oct 27, 1992
Days to decision	47 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Mitek Surgical Products, Inc.</b>
Location	Dedham, MA, US
Contact	ROBERT P ZOLETTI
510(k) history	31 submissions · 26 cleared · 1989-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k924665/> 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 14, 2026