

**K930893 MITEK SUPER QUICKANCHOR, MODIFICATION**Dec 8, 1993  
313 days to decisionK930893 · Product code: **JDR** · Orthopedic  
Source: <https://www.510kdatabase.net/k930893/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Staple, Fixation, Bone (JDR)
Date received	Jan 29, 1993
Decision date	Dec 8, 1993
Days to decision	313 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k930893/> 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