

**K032078 MITEK MICRO QUICKANCHOR**Aug 25, 2003  
49 days to decisionK032078 · Product code: **NOV** · Orthopedic  
Source: <https://www.510kdatabase.net/k032078/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Anchor, Suture, Bone Fixation, Metallic (NOV)
Date received	Jul 7, 2003
Decision date	Aug 25, 2003
Days to decision	49 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Mitek Worldwide</b>
Location	Norwood, MA, US
Contact	SERGIO J GADALETA
510(k) history	10 submissions · 10 cleared · 2002-2004

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k032078/> 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