

**K052631 SUPER QUICKANCHOR PLUS (WITH ORTHOCORD SUTURE)**Oct 21, 2005  
25 days to decisionK052631 · Product code: JDR · Orthopedic  
Source: <https://www.510kdatabase.net/k052631/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Staple, Fixation, Bone (JDR)
Date received	Sep 26, 2005
Decision date	Oct 21, 2005
Days to decision	25 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Depuy Mitek, A Johnson &amp; Johnson Company</b>
Location	Norwood, MA, US
Contact	DENISE LUCIANO
510(k) history	58 submissions · 58 cleared · 2004-2015

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k052631/> Data sourced from FDA 510(k) public records (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. - Generated May 14, 2026