

**K051989 GII QUICKANCHOR PLUS WITH ORTHOCORD
SUTURE**Aug 5, 2005
14 days to decisionK051989 · Product code: JDR · Orthopedic
Source: <https://www.510kdatabase.net/k051989/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Staple, Fixation, Bone (JDR)
Date received	Jul 22, 2005
Decision date	Aug 5, 2005
Days to decision	14 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k051989/> 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