

**K061801 2.8 & 3.5MM LACTOSCREW SUTURE ANCHORS**Aug 7, 2006  
42 days to decisionK061801 · Product code: **JDR** · Orthopedic  
Source: <https://www.510kdatabase.net/k061801/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Staple, Fixation, Bone (JDR)
Date received	Jun 26, 2006
Decision date	Aug 7, 2006
Days to decision	42 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Arthrotek, Inc.</b>
Location	West Hartford, CT, US
Contact	SUSAN ALEXANDER
510(k) history	17 submissions · 17 cleared · 1987-2006

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k061801/> 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