

**K061465 KMI CEMENT RESTRICTOR IMPLANT**Oct 5, 2006  
132 days to decisionK061465 · Product code: **JDK** · Orthopedic  
Source: <https://www.510kdatabase.net/k061465/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - U
Submission type	Traditional
Device classification	Prosthesis, Hip, Cement Restrictor (JDK)
Date received	May 26, 2006
Decision date	Oct 5, 2006
Days to decision	132 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Kinetikos Medical, Inc.</b>
Location	San Diego, CA, US
Contact	JOHN G SPAMPINATO
510(k) history	19 submissions · 16 cleared · 1994-2006

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k061465/> 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 19, 2026