

**K051371 INTERBODY INNOVATIONS CEMENT RESTRICTOR**Dec 6, 2005  
194 days to decisionK051371 · Product code: **JDK** · Orthopedic  
Source: <https://www.510kdatabase.net/k051371/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - U
Submission type	Traditional
Device classification	Prosthesis, Hip, Cement Restrictor (JDK)
Date received	May 26, 2005
Decision date	Dec 6, 2005
Days to decision	194 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Interbody Innovations, Llp</b>
Location	Clearwater, FL, US
Contact	IAN GORDON
510(k) history	4 submissions · 1 cleared · 2005-2008

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k051371/> 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