

**K021788 NBD CEMENT RESTRICTOR DEVICE**Oct 4, 2002  
126 days to decisionK021788 · Product code: **JDK** · Orthopedic  
Source: <https://www.510kdatabase.net/k021788/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - U
Submission type	Traditional
Device classification	Prosthesis, Hip, Cement Restrictor (JDK)
Date received	May 31, 2002
Decision date	Oct 4, 2002
Days to decision	126 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>New Business Development, LLC</b>
Location	Woodstock, GA, US
Contact	TIM B LUSBY
510(k) history	4 submissions · 3 cleared · 2002-2007

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k021788/> 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