

**K053080 GRAFTCAGE ACX**Dec 1, 2005  
29 days to decisionK053080 · Product code: **MQP** · Orthopedic  
Source: <https://www.510kdatabase.net/k053080/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spinal Vertebral Body Replacement Device (MQP)
Date received	Nov 2, 2005
Decision date	Dec 1, 2005
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Osteotech, Inc.</b>
Location	San Mateo, CA, US
Contact	CHRISTOPHER TALBOT
510(k) history	24 submissions · 21 cleared · 1985-2008

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k053080/> 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