

**K051781 GRAFTCAGE TLX**Dec 16, 2005  
168 days to decisionK051781 · Product code: **MQP** · Orthopedic  
Source: <https://www.510kdatabase.net/k051781/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spinal Vertebral Body Replacement Device (MQP)
Date received	Jul 1, 2005
Decision date	Dec 16, 2005
Days to decision	168 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k051781/> 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