

**K062666 VUSION TS, PS, AND CS PARTIAL VERTEBRAL BODY REPLACEMENTS**Jan 25, 2007  
140 days to decisionK062666 · Product code: **MQP** · Orthopedic  
Source: <https://www.510kdatabase.net/k062666/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spinal Vertebral Body Replacement Device (MQP)
Date received	Sep 7, 2006
Decision date	Jan 25, 2007
Days to decision	140 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Ortho Development Corp.</b>
Location	Draper, UT, US
Contact	WILLIAM J GRIFFIN
510(k) history	45 submissions · 43 cleared · 1996-2025

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k062666/> Data sourced from FDA 510(k) public records (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. - Generated May 19, 2026