

**K090899 DEPUY SPINE LATERAL SYSTEM**May 19, 2009  
48 days to decisionK090899 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k090899/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Apr 1, 2009
Decision date	May 19, 2009
Days to decision	48 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Depuy Spine, Inc.</b>
Location	Raynham, MA, US
Contact	DENISE DUCHENE
510(k) history	68 submissions · 67 cleared · 2004-2020

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k090899/> 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 June 19, 2026