

**K080411 PLATEAU SPACER SYSTEM**May 15, 2008  
90 days to decisionK080411 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k080411/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Feb 15, 2008
Decision date	May 15, 2008
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Life Spine</b>
Location	Hoffman Estates, IL, US
Contact	REBECCA M BROOKS
510(k) history	36 submissions · 34 cleared · 2006-2018

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k080411/> 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