

**K133717 LONGBOW SPACER SYSTEM**Sep 19, 2014  
288 days to decisionK133717 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k133717/>**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	Dec 5, 2013
Decision date	Sep 19, 2014
Days to decision	288 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Life Spine</b>
Location	Hoffman Estates, IL, US
Contact	RANDY LEWIS
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/k133717/> 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