

K131077 LIFE SPINE PLATEAU SPACER SYSTEMJul 17, 2013
91 days to decisionK131077 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k131077/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Apr 17, 2013
Decision date	Jul 17, 2013
Days to decision	91 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Life Spine
Location	Hoffman Estates, IL, US
Contact	RANDY LEWIS
510(k) history	36 submissions · 34 cleared · 2006-2018

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k131077/> 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 June 19, 2026