

**K223430 ProLift Micro Expandable Spacer System**May 4, 2023  
171 days to decisionK223430 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k223430/>**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	Nov 14, 2022
Decision date	May 4, 2023
Days to decision	171 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Life Spine, Inc.</b>
Location	Hoffman Estates, IL, US
Contact	Angela Batker
510(k) history	82 submissions · 82 cleared · 2011-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k223430/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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