

K251502 TruLift® Lateral Expandable Spacer & Lateral Buttress Plate System

Jul 7, 2025
53 days to decision

K251502 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k251502/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	May 15, 2025
Decision date	Jul 7, 2025
Days to decision	53 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k251502/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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