FDA 510(k) Premarket Notification

K250373 ProLift Expandable Spacer System

CLEARED

Apr 9, 2025
58 days to decision

K250373 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k250373/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Feb 10, 2025
Decision date	Apr 9, 2025
Days to decision	58 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