FDA 510(k) Premarket Notification

K212520 ProLift Micro Expandable Spacer System

CLEARED

Oct 8, 2021
59 days to decision

K212520 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k212520/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Aug 10, 2021
Decision date	Oct 8, 2021
Days to decision	59 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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k212520/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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