FDA 510(k) Premarket Notification

K242826 ProLift Wedge Expandable Spacer System

CLEARED

Oct 8, 2024
20 days to decision

K242826 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k242826/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Sep 18, 2024
Decision date	Oct 8, 2024
Days to decision	20 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