

K190488 ProLift Expandable SystemJul 23, 2019
145 days to decisionK190488 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k190488/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Feb 28, 2019
Decision date	Jul 23, 2019
Days to decision	145 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Life Spine, Inc.
Location	Hoffman Estates, IL, US
Contact	Angela Batker
Website	http://www.lifespine.com/
510(k) history	82 submissions · 82 cleared · 2011-2026

Life Spine, Inc. is a spinal medical device company headquartered in Huntley, Illinois. Founded in 2004, the company develops innovative solutions for spinal pathology across the cervical, thoracic, and lumbar spine. Life Spine serves 32 countries and employs over 70 people worldwide. The company has received FDA 510(k) clearances from total submissions since 2011. Life Spine specializes exclusively in Orthopedic devices, with a focus on minimally invasive spinal fusion solutions. The latest clearance was in 2026, confirming active regulatory engagement. Life Spine's prod...
