FDA 510(k) Premarket Notification

K092815 S-LIF INTERVERTEBRAL BODY FUSION DEVICE

CLEARED

Jun 30, 2010
289 days to decision

K092815 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k092815/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Sep 14, 2009
Decision date	Jun 30, 2010
Days to decision	289 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Spinefrontier, Inc.**
Location	Beverly, MA, US
Contact	JOHN SULLIVAN
510(k) history	24 submissions · 24 cleared · 2007-2020