FDA 510(k) Premarket Notification

K071724 LUCENT

CLEARED

Nov 9, 2007
137 days to decision

K071724 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k071724/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Jun 25, 2007
Decision date	Nov 9, 2007
Days to decision	137 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Spinal Elements, Inc.**
Location	Carlsbad, CA, US
Contact	KERRI DIMARTINO
Website	https://www.spinalelements.com
510(k) history	48 submissions · 48 cleared · 2007-2026