FDA 510(k) Premarket Notification

K090566 PATHWAY AVID

CLEARED

Jun 1, 2009
90 days to decision

K090566 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k090566/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Mar 3, 2009
Decision date	Jun 1, 2009
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Custom Spine, Inc.**
Location	Conway, NH, US
Contact	SAAD ATTIYAH
510(k) history	12 submissions · 12 cleared · 2005-2015