FDA 510(k) Premarket Notification

K090899 DEPUY SPINE LATERAL SYSTEM

CLEARED

May 19, 2009
48 days to decision

K090899 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k090899/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Apr 1, 2009
Decision date	May 19, 2009
Days to decision	48 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Depuy Spine, Inc.**
Location	Raynham, MA, US
Contact	DENISE DUCHENE
510(k) history	68 submissions · 67 cleared · 2004-2020