FDA 510(k) Premarket Notification

K090376 SPINESMITH CYNCH SPINAL SYSTEM

CLEARED

Apr 1, 2009
43 days to decision

K090376 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k090376/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Feb 17, 2009
Decision date	Apr 1, 2009
Days to decision	43 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Spine Smith Partners L.P.**
Location	Austin, TX, US
Contact	ROBERT JONES
510(k) history	9 submissions · 9 cleared · 2008-2012