FDA 510(k) Premarket Notification

K131540 INTERFUSE L INTERVERTEBRAL BODY FUSION DEVICE

CLEARED

Sep 3, 2013
97 days to decision

K131540 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k131540/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	May 29, 2013
Decision date	Sep 3, 2013
Days to decision	97 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Vertebral Technologies, Inc.**
Location	Andover, MN, US
Contact	SURESH GHAI
510(k) history	7 submissions · 7 cleared · 2008-2013