FDA 510(k) Premarket Notification

CLEARED

K091988 MODIFICATION TO: INTERFUSE INTERVERTEBRAL BODY FUSION DEVICE , MODEL 9076

Jul 30, 2009
28 days to decision

K091988 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k091988/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Jul 2, 2009
Decision date	Jul 30, 2009
Days to decision	28 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