FDA 510(k) Premarket Notification

CLEARED

K093675 INTERFUSE INVERTEBRAL BODY FUSION DEVICE MODEL 9076

Dec 23, 2009
26 days to decision

K093675 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k093675/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Nov 27, 2009
Decision date	Dec 23, 2009
Days to decision	26 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