FDA 510(k) Premarket Notification

K110226 INTERFUSE T INVERTEBRAL BODY FUSION DEVICE

CLEARED

May 27, 2011
122 days to decision

K110226 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k110226/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Jan 25, 2011
Decision date	May 27, 2011
Days to decision	122 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