FDA 510(k) Premarket Notification

CLEARED

K080673 INTERFUSE INTERVERTEBRAL BODY FUSION DEVICE

Jun 10, 2008
92 days to decision

K080673 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k080673/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Mar 10, 2008
Decision date	Jun 10, 2008
Days to decision	92 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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