FDA 510(k) Premarket Notification

K160874 Nanovis Intervertebral Body Fusion System and FortiCore®

CLEARED

Jun 28, 2016
90 days to decision

K160874 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k160874/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Mar 30, 2016
Decision date	Jun 28, 2016
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Nanovis, LLC**
Location	San Diego, CA, US
Contact	Matthew Hedrick
510(k) history	10 submissions · 10 cleared · 2011-2024