FDA 510(k) Premarket Notification

K142594 nva, nvp, nvt

CLEARED

Nov 28, 2014
74 days to decision

K142594 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k142594/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Sep 15, 2014
Decision date	Nov 28, 2014
Days to decision	74 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Nvision Biomedical Technologies, LLC**
Location	San Antonio, TN, US
Contact	Brian Kieser
510(k) history	10 submissions · 10 cleared · 2014-2019