

**K133721 INFILL 41-TLIF CONVEX OBLIQUE DEVICE, INFILL
43-TLIF CONTOUR OBLIQUE, INFILL 44-TLIF CONTOUR
OBLIQUE AND THE INFILL 60**Mar 31, 2014
116 days to decisionK133721 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k133721/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Dec 5, 2013
Decision date	Mar 31, 2014
Days to decision	116 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Pinnacle Spine Group, LLC
Location	Dallas, TX, US
Contact	Rebecca K Pine
510(k) history	12 submissions · 12 cleared · 2011-2017

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k133721/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 16, 2026