

K113593 NEXTGEN ALTIUS OCT SYSTEMApr 17, 2012
134 days to decisionK113593 · Product code: **KWP** · Orthopedic
Source: <https://www.510kdatabase.net/k113593/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Interlaminar (KWP)
Date received	Dec 5, 2011
Decision date	Apr 17, 2012
Days to decision	134 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Biomet Spine (Aka Ebi, LLC)
Location	Parsippany, NJ, US
Contact	VIVIAN KELLY
510(k) history	13 submissions · 13 cleared · 2010-2014

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k113593/> 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 17, 2026