

K133363 INTERBRIDGE INTERSPINOUS POSTERIOR FIXATION SYSTEMMar 7, 2014
126 days to decisionK133363 · Product code: **PEK** · Orthopedic
Source: <https://www.510kdatabase.net/k133363/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spinous Process Plate (PEK)
Date received	Nov 1, 2013
Decision date	Mar 7, 2014
Days to decision	126 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Ldr Spine USA, Inc.
Location	Austin, TX, US
Contact	MEREDITH MAY
510(k) history	10 submissions · 10 cleared · 2011-2017

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k133363/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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