

**K070341 LDR SPINE EASYSPINE POSTERIOR
OSTEOSYNTHESIS SYSTEM**Aug 8, 2007
183 days to decisionK070341 · Product code: **KWP** · Orthopedic
Source: <https://www.510kdatabase.net/k070341/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Interlaminar (KWP)
Date received	Feb 6, 2007
Decision date	Aug 8, 2007
Days to decision	183 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Ldr Spine USA
Location	Austin, TX, US
Contact	JAMES BURROWS
510(k) history	25 submissions · 25 cleared · 2005-2016

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k070341/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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