

K120760 LDR SPINE SPINETUNE TL SPINAL SYSTEMJun 27, 2012
106 days to decisionK120760 · Product code: **MNH** · Orthopedic
Source: <https://www.510kdatabase.net/k120760/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Orthosis, Spondylolisthesis Spinal Fixation (MNH)
Date received	Mar 13, 2012
Decision date	Jun 27, 2012
Days to decision	106 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k120760/> 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