

**K102265 LDR SPINE USA C-PLATE, TL ANTERIOR CERVICAL PLATE SYSTEM**

Sep 29, 2010  
50 days to decision

K102265 · Product code: **KWQ** · Orthopedic  
Source: <https://www.510kdatabase.net/k102265/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Aug 10, 2010
Decision date	Sep 29, 2010
Days to decision	50 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Ldr Spine USA</b>
Location	Austin, TX, US
Contact	BECKINAM NOWATZKE
510(k) history	25 submissions · 25 cleared · 2005-2016

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k102265/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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