FDA 510(k) Premarket Notification

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

CLEARED

Sep 29, 2010
50 days to decision

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

SUBMISSION DETAILS

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

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