FDA 510(k) Premarket Notification

K070341 LDR SPINE EASYSPINE POSTERIOR OSTEOSYNTHESIS SYSTEM

CLEARED

Aug 8, 2007
183 days to decision

K070341 · Product code: **KWP** · Orthopedic
Source: https://www.510kdatabase.net/k070341/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Interlaminal (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