FDA 510(k) Premarket Notification

K070922 MODIFICATION TO SPINAL USA VBR SYSTEM

CLEARED

Sep 19, 2007
170 days to decision

K070922 · Product code: **KWQ** · Orthopedic
Source: https://www.510kdatabase.net/k070922/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Apr 2, 2007
Decision date	Sep 19, 2007
Days to decision	170 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Spinal USA**
Location	Brandon, MS, US
Contact	JEFFREY JOHNSON
510(k) history	23 submissions · 23 cleared · 2006-2013