FDA 510(k) Premarket Notification

K051716 VERTEBRON PSS PEDICLE SCREW SYSTEM

CLEARED

Sep 7, 2005
72 days to decision

K051716 · Product code: **KWP** · Orthopedic
Source: https://www.510kdatabase.net/k051716/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Spinal Interlaminal (KWP)
Date received	Jun 27, 2005
Decision date	Sep 7, 2005
Days to decision	72 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Vertebron, Inc.**
Location	Stratford, CT, US
Contact	LUIS NESPRIDO
510(k) history	11 submissions · 11 cleared · 2004-2008