

K061752 SPINAL USA VBR SYSTEM

Aug 15, 2006
55 days to decision

K061752 · Product code: **MQP** · Orthopedic
Source: <https://www.510kdatabase.net/k061752/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spinal Vertebral Body Replacement Device (MQP)
Date received	Jun 21, 2006
Decision date	Aug 15, 2006
Days to decision	55 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

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Device record: <https://www.510kdatabase.net/k061752/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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