FDA 510(k) Premarket Notification

K132029 VAULT-C INTERVERTEBRAL BODY FUSION DEVICE

CLEARED

Nov 25, 2013
147 days to decision

K132029 · Product code: **OVE** · Orthopedic
Source: https://www.510kdatabase.net/k132029/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Integrated Fixation, Cervical (OVE)
Date received	Jul 1, 2013
Decision date	Nov 25, 2013
Days to decision	147 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k132029/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 17, 2026