FDA 510(k) Premarket Notification

K061353 TRINICA ANTERIOR LUMBAR PLATE SYSTEM

CLEARED

Aug 30, 2006
107 days to decision

K061353 · Product code: **KWQ** · Orthopedic
Source: https://www.510kdatabase.net/k061353/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	May 15, 2006
Decision date	Aug 30, 2006
Days to decision	107 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Zimmer Spine, Inc.**
Location	Minneapolis, MN, US
Contact	TIM CRABTREE
510(k) history	38 submissions · 35 cleared · 2004-2016

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k061353/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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