FDA 510(k) Premarket Notification

K071880 Optima ZS Transition Screw

CLEARED

Oct 5, 2007
88 days to decision

K071880 · Product code: **MNI** · Orthopedic
Source: https://www.510kdatabase.net/k071880/

SUBMISSION DETAILS

Decision	Substantially Equivalent - U
Submission type	Traditional
Device classification	Orthosis, Spinal Pedicle Fixation (MNI)
Date received	Jul 9, 2007
Decision date	Oct 5, 2007
Days to decision	88 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k071880/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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