

**K081597 MODIFICATION TO VERTEBRON PSS PEDICLE
SCREW SYSTEM**Jul 2, 2008
26 days to decisionK081597 · Product code: **MNI** · Orthopedic
Source: <https://www.510kdatabase.net/k081597/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Orthosis, Spinal Pedicle Fixation (MNI)
Date received	Jun 6, 2008
Decision date	Jul 2, 2008
Days to decision	26 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k081597/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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