

**K070295 SINGLE PLANAR MULTI AXIAL (SPMA) PEDICLE SCREW SYSTEM**May 21, 2007  
110 days to decisionK070295 · Product code: MNI · Orthopedic  
Source: <https://www.510kdatabase.net/k070295/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthosis, Spinal Pedicle Fixation (MNI)
Date received	Jan 31, 2007
Decision date	May 21, 2007
Days to decision	110 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Trinity Orthopedics, LLC</b>
Location	San Diego,, CA, US
Contact	FLOYD G LARSON
510(k) history	2 submissions · 2 cleared · 2007-2008

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k070295/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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