

**K992369 SYNERGY INTEGRAL SCREWS, MODELS, 2161,2162,2163,2164,2165,2166,2167,2168,2169,2210,2211,2212,2215,2216,2217,2220,2221,22**

Oct 13, 1999  
90 days to decision

K992369 · Product code: **MNI** · Orthopedic  
Source: <https://www.510kdatabase.net/k992369/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthosis, Spinal Pedicle Fixation (MNI)
Date received	Jul 15, 1999
Decision date	Oct 13, 1999
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Interpore Cross Intl.</b>
Location	Irvine, CA, US
Contact	LYNN M RODARTI
510(k) history	39 submissions · 38 cleared · 1998-2005

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k992369/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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