

**K041407 TITANIUM MESH IMPLANT**Dec 29, 2004  
216 days to decisionK041407 · Product code: **MQP** · Orthopedic  
Source: <https://www.510kdatabase.net/k041407/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spinal Vertebral Body Replacement Device (MQP)
Date received	May 27, 2004
Decision date	Dec 29, 2004
Days to decision	216 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Interpore Cross Intl.</b>
Location	Irvine, CA, US
Contact	WENDY SPIELBERGER
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/k041407/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 20, 2026