

**K032371 TITANIUM MESH SYSTEM**Feb 12, 2004  
195 days to decisionK032371 · Product code: **MQP** · Orthopedic  
Source: <https://www.510kdatabase.net/k032371/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spinal Vertebral Body Replacement Device (MQP)
Date received	Aug 1, 2003
Decision date	Feb 12, 2004
Days to decision	195 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Encore Medical, L.P.</b>
Location	Austin, TX, US
Contact	KIMBERLY L PRUITT
510(k) history	81 submissions · 81 cleared · 2002-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k032371/> 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 14, 2026