

**K092336 C-THRU SPINAL SYSTEM**Oct 15, 2009  
72 days to decisionK092336 · Product code: **MQP** · Orthopedic  
Source: <https://www.510kdatabase.net/k092336/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spinal Vertebral Body Replacement Device (MQP)
Date received	Aug 4, 2009
Decision date	Oct 15, 2009
Days to decision	72 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Ebi, L.P.</b>
Location	Parsippany, NJ, US
Contact	VIVIAN KELLY
510(k) history	95 submissions · 94 cleared · 1997-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k092336/> 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 18, 2026