

**K073440 FOUNDATION CAGE**Apr 24, 2008  
139 days to decisionK073440 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k073440/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Dec 7, 2007
Decision date	Apr 24, 2008
Days to decision	139 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Corelink, LLC</b>
Location	Round Rock, TX, US
Contact	J.D. WEBB
510(k) history	35 submissions · 35 cleared · 2008-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k073440/> 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