

**K150847 Foundation Interbody Devices**Aug 13, 2015  
135 days to decisionK150847 · Product code: **ODP** · Orthopedic  
Source: <https://www.510kdatabase.net/k150847/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Cervical (ODP)
Date received	Mar 31, 2015
Decision date	Aug 13, 2015
Days to decision	135 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Corelink, LLC</b>
Location	Round Rock, TX, US
Contact	Sharon Gregory
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/k150847/> 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 17, 2026