

**K113540 RELIANCE LUMBAR IBF SYSTEM**Mar 12, 2012  
103 days to decisionK113540 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k113540/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Nov 30, 2011
Decision date	Mar 12, 2012
Days to decision	103 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Reliance Medical Systems, LLC</b>
Location	Sandy, UT, US
Contact	Bret M Berry
510(k) history	24 submissions · 24 cleared · 2007-2022

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k113540/> 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 16, 2026