

**K963812 LM BONE ANCHOR (ORTHOPEDICS)**Nov 12, 1996  
64 days to decisionK963812 · Product code: **MBI** · Orthopedic  
Source: <https://www.510kdatabase.net/k963812/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Fastener, Fixation, Nondegradable, Soft Tissue (MBI)
Date received	Sep 9, 1996
Decision date	Nov 12, 1996
Days to decision	64 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Li Medical Technologies, Inc.</b>
Location	Greenwich, CT, US
Contact	RHODEMAN LI
510(k) history	17 submissions · 17 cleared · 1993-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k963812/> 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