

**K933653 MS 2500**Mar 23, 1994  
238 days to decisionK933653 · Product code: **IPF** · Physical MedicineSource: <https://www.510kdatabase.net/k933653/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Muscle, Powered (IPF)
Date received	Jul 28, 1993
Decision date	Mar 23, 1994
Days to decision	238 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Lsi Intl., Inc.</b>
Location	Overland Park, KS, US
Contact	STEVE PAPA
510(k) history	5 submissions · 5 cleared · 1994-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k933653/> 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 June 29, 2026