

**K903429 LIDO LINEA**Aug 14, 1990  
14 days to decisionK903429 · Product code: **IKK** · Physical Medicine  
Source: <https://www.510kdatabase.net/k903429/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Isokinetic Testing And Evaluation (IKK)
Date received	Jul 31, 1990
Decision date	Aug 14, 1990
Days to decision	14 days
Third-party review	No

**APPLICANT**

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Company	<b>Loredan Biomedical, Inc.</b>
Location	Mchenry, IL, US
Contact	KAREN COMISSO
510(k) history	9 submissions · 9 cleared · 1983-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k903429/>; 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 30, 2026