

**K880017 LIDO(TM) LIFT**Feb 26, 1988  
52 days to decisionK880017 · Product code: **IKK** · Physical MedicineSource: <https://www.510kdatabase.net/k880017/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Isokinetic Testing And Evaluation (IKK)
Date received	Jan 5, 1988
Decision date	Feb 26, 1988
Days to decision	52 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/k880017/>, 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