

**K925387 LIBERATOR**Mar 29, 1993  
154 days to decisionK925387 · Product code: **ILQ** · Physical Medicine  
Source: <https://www.510kdatabase.net/k925387/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Communication, Powered (ILQ)
Date received	Oct 26, 1992
Decision date	Mar 29, 1993
Days to decision	154 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Prentke Romich Co.</b>
Location	Wooster, OH, US
Contact	MICHAEL MOLTER
510(k) history	9 submissions · 7 cleared · 1990-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k925387/> 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