

**K853303 LIFELINC LEADWIRE**Sep 18, 1985  
43 days to decisionK853303 · Product code: **BZQ** · Anesthesiology  
Source: <https://www.510kdatabase.net/k853303/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Breathing Frequency (BZQ)
Date received	Aug 6, 1985
Decision date	Sep 18, 1985
Days to decision	43 days
Third-party review	No

**APPLICANT**

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Company	<b>Tronomed, Inc.</b>
Location	Mchenry, IL, US
Contact	JIM LELTEN
510(k) history	12 submissions · 12 cleared · 1977-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k853303/>; 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 27, 2026