

**K896300 LDW SPIROMETRY SYSTEM**Apr 3, 1990  
155 days to decisionK896300 · Product code: **BZG** · Anesthesiology  
Source: <https://www.510kdatabase.net/k896300/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spirometer, Diagnostic (BZG)
Date received	Oct 30, 1989
Decision date	Apr 3, 1990
Days to decision	155 days
Third-party review	No

**APPLICANT**

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Company	<b>Lanpher Diagnostic Workstations, Inc.</b>
Location	Atherton, CA, US
Contact	TED W LANPHER
510(k) history	1 submissions · 1 cleared · 1990-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k896300/>; 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 26, 2026