FDA 510(k) Premarket Notification

# K896300 LDW SPIROMETRY SYSTEM

**CLEARED**

Apr 3, 1990
155 days to decision

K896300 · 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

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