

**K013251 SPIRIT 300**Dec 18, 2001  
81 days to decisionK013251 · Product code: **BYJ** · Anesthesiology  
Source: <https://www.510kdatabase.net/k013251/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Liquid-oxygen, Portable (BYJ)
Date received	Sep 28, 2001
Decision date	Dec 18, 2001
Days to decision	81 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Caire, Inc.</b>
Location	Littleton, CO, US
Contact	ROGER BRIESE
510(k) history	7 submissions · 7 cleared · 1995-2025

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k013251/> 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