

**K896121 SCHILLER MODEL SP-200 SPIROVIT(TM)**Jan 9, 1990  
77 days to decisionK896121 · Product code: **BZG** · Anesthesiology  
Source: <https://www.510kdatabase.net/k896121/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spirometer, Diagnostic (BZG)
Date received	Oct 24, 1989
Decision date	Jan 9, 1990
Days to decision	77 days
Third-party review	No

**APPLICANT**

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Company	<b>Schiller America, Inc.</b>
Location	Tustin, CA, US
Contact	GODFREY J FLETCHER
510(k) history	10 submissions · 10 cleared · 1990-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k896121/>, 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