

**K832401 RESPIROMETER**Apr 30, 1984  
285 days to decisionK832401 · Product code: **BZG** · AnesthesiologySource: <https://www.510kdatabase.net/k832401/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spirometer, Diagnostic (BZG)
Date received	Jul 20, 1983
Decision date	Apr 30, 1984
Days to decision	285 days
Third-party review	No

**APPLICANT**

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Company	<b>Kinetix</b>
Location	Walker, MI, US
510(k) history	4 submissions · 4 cleared · 1983-1984

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k832401/> 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