

**K213872 CompAS2**Jul 15, 2022  
214 days to decisionK213872 · Product code: **BZG** · Anesthesiology  
Source: <https://www.510kdatabase.net/k213872/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spirometer, Diagnostic (BZG)
Date received	Dec 13, 2021
Decision date	Jul 15, 2022
Days to decision	214 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Morgan Scientific, Inc.</b>
Location	Haverhill, MA, US
Contact	Deborah Cornish
510(k) history	6 submissions · 6 cleared · 2002-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k213872/> 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