

**K083199 CHARTIS CONSOLE**Jun 25, 2009  
238 days to decisionK083199 · Product code: **BZG** · Anesthesiology  
Source: <https://www.510kdatabase.net/k083199/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spirometer, Diagnostic (BZG)
Date received	Oct 30, 2008
Decision date	Jun 25, 2009
Days to decision	238 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Pulmonx, Inc.</b>
Location	Palo Alto, CA, US
Contact	SUZON LOMMEL
510(k) history	4 submissions · 4 cleared · 2009-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k083199/> 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 24, 2026