

**K062689 SIDESTREAM PLUS**Feb 16, 2007  
158 days to decisionK062689 · Product code: **CAF** · Anesthesiology  
Source: <https://www.510kdatabase.net/k062689/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Nebulizer (direct Patient Interface) (CAF)
Date received	Sep 11, 2006
Decision date	Feb 16, 2007
Days to decision	158 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Respironics , Ltd.</b>
Location	Cedar Grove, NJ, US
Contact	Lauren Ziegler
510(k) history	1 submissions · 1 cleared · 2007-2007

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k062689/> 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