

**K830895 SINGLE USE NEBULIZER, ADJUST. AIR-**Apr 28, 1983  
38 days to decisionK830895 · Product code: **CAF** · AnesthesiologySource: <https://www.510kdatabase.net/k830895/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Nebulizer (direct Patient Interface) (CAF)
Date received	Mar 21, 1983
Decision date	Apr 28, 1983
Days to decision	38 days
Third-party review	No

**APPLICANT**

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Company	<b>Airlife, Inc.</b>
Location	Mchenry, IL, US
510(k) history	76 submissions · 76 cleared · 1979-1984

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k830895/> 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