

**K810947 AIR ENTRAINMENT TRACH T**Jun 2, 1981  
56 days to decisionK810947 · Product code: **BYF** · Anesthesiology  
Source: <https://www.510kdatabase.net/k810947/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mask, Oxygen, Low Concentration, Venturi (BYF)
Date received	Apr 7, 1981
Decision date	Jun 2, 1981
Days to decision	56 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/k810947/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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