

**K810335 RESPIRIZER**Mar 20, 1981  
39 days to decisionK810335 · Product code: **KFZ** · Anesthesiology  
Source: <https://www.510kdatabase.net/k810335/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Humidifier, Non-direct Patient Interface (home-use) (KFZ)
Date received	Feb 9, 1981
Decision date	Mar 20, 1981
Days to decision	39 days
Third-party review	No

**APPLICANT**

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Company	<b>Respirizer Co., Div. Campillary Systems, Inc.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1981-1981

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k810335/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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