

**K013111 EXHALE PROBE**Oct 1, 2001  
13 days to decisionK013111 · Product code: **EOQ** · Radiology  
Source: <https://www.510kdatabase.net/k013111/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bronchoscope (flexible Or Rigid) (EOQ)
Date received	Sep 18, 2001
Decision date	Oct 1, 2001
Days to decision	13 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Broncus Technologies, Inc.</b>
Location	Mountain View, CA, US
Contact	TIMOTHY R WILLIAMS
510(k) history	11 submissions · 11 cleared · 1998-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k013111/> 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 17, 2026