

**K954707 BRONCHOSCOPE**Nov 22, 1995  
41 days to decisionK954707 · Product code: **EOQ** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k954707/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bronchoscope (flexible Or Rigid) (EOQ)
Date received	Oct 12, 1995
Decision date	Nov 22, 1995
Days to decision	41 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Fujinon, Inc.</b>
Location	Washington, DC, US
Contact	GARY A ADLER
510(k) history	30 submissions · 30 cleared · 1990-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k954707/> 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