

**K832010 BRONCHO-FIBERSCOPE**Aug 1, 1983  
39 days to decisionK832010 · Product code: **EOQ** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k832010/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bronchoscope (flexible Or Rigid) (EOQ)
Date received	Jun 23, 1983
Decision date	Aug 1, 1983
Days to decision	39 days
Third-party review	No

**APPLICANT**

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Company	<b>KARL STORZ Endoscopy-America, Inc.</b>
Location	Mchenry, IL, US
510(k) history	361 submissions · 361 cleared · 1980-2024

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

Device record: <https://www.510kdatabase.net/k832010/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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