

**K810007 PENTAX UPPER G.I. FIBERSCOPE, #FG-34**Jan 15, 1981  
10 days to decisionK810007 · Product code: **GDB** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k810007/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Endoscope, Fiber Optic (GDB)
Date received	Jan 5, 1981
Decision date	Jan 15, 1981
Days to decision	10 days
Third-party review	No

**APPLICANT**

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Company	<b>Pentax Precision Instrument Corp.</b>
Location	Mchenry, IL, US
510(k) history	67 submissions · 67 cleared · 1979-2004

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k810007/> 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 18, 2026