

K832006 PENTAX FD-34A DUODENOFIBERSCOPEAug 1, 1983
39 days to decisionK832006 · Product code: **FDT** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k832006/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Duodenoscope And Accessories, Flexible/rigid (FDT)
Date received	Jun 23, 1983
Decision date	Aug 1, 1983
Days to decision	39 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k832006/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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