

K820239 RETRIEVAL FORCEPSMar 26, 1982
56 days to decisionK820239 · Product code: **KOD** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k820239/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Urological (KOD)
Date received	Jan 29, 1982
Decision date	Mar 26, 1982
Days to decision	56 days
Third-party review	No

APPLICANT

Company	American Endoscopy, Inc.
Location	Mchenry, IL, US
510(k) history	19 submissions · 19 cleared · 1982-1985

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Device record: <https://www.510kdatabase.net/k820239/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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