

K820431 E.R.C.P. CANNULASApr 9, 1982
51 days to decisionK820431 · Product code: **ODD** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k820431/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Endoscopic Retrograde Cholangiopancreatography (ercp) Cannula (ODD)
Date received	Feb 17, 1982
Decision date	Apr 9, 1982
Days to decision	51 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/k820431/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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