

K833417 ENDOSCOPIC RETROGRADENov 28, 1983
56 days to decisionK833417 · Product code: **ODD** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k833417/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Endoscopic Retrograde Cholangiopancreatography (ercp) Cannula (ODD)
Date received	Oct 3, 1983
Decision date	Nov 28, 1983
Days to decision	56 days
Third-party review	No

APPLICANT

Company	Microvasive
Location	Mchenry, IL, US
510(k) history	18 submissions · 18 cleared · 1983-1987

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k833417/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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