

**K833920 KELLY CHOLANGIOCATH**Dec 27, 1983  
43 days to decisionK833920 · Product code: **FGE** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k833920/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stents, Drains And Dilators For The Biliary Ducts (FGE)
Date received	Nov 14, 1983
Decision date	Dec 27, 1983
Days to decision	43 days
Third-party review	No

**APPLICANT**

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Company	<b>Perry, Div.</b>
Location	Mchenry, IL, US
510(k) history	6 submissions · 6 cleared · 1983-1985

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k833920/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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