

**K905714 ERCP CANNULA**Feb 1, 1991  
42 days to decisionK905714 · Product code: **FGE** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k905714/>**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	Dec 21, 1990
Decision date	Feb 1, 1991
Days to decision	42 days
Third-party review	No

**APPLICANT**

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Company	<b>Everest Medical Corp.</b>
Location	Brooklyn Center, MN, US
Contact	DAVID PARINS
510(k) history	21 submissions · 21 cleared · 1987-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k905714/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 29, 2026