FDA 510(k) Premarket Notification

K905714 ERCP CANNULA

CLEARED

Feb 1, 1991
42 days to decision

K905714 · Product code: **FGE** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k905714/

SUBMISSION DETAILS

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

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