FDA 510(k) Premarket Notification

K833920 KELLY CHOLANGIOCATH

CLEARED

Dec 27, 1983
43 days to decision

K833920 · Product code: **FGE** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k833920/

SUBMISSION DETAILS

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

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