FDA 510(k) Premarket Notification

K855142 ANGIOMED PERCUTANEOUS TRANSHEPATIC CHOLANGIOGTAPHY

CLEARED

Mar 11, 1986
75 days to decision

K855142 · Product code: **FGE** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k855142/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stents, Drains And Dilators For The Biliary Ducts (FGE)
Date received	Dec 26, 1985
Decision date	Mar 11, 1986
Days to decision	75 days
Third-party review	No

APPLICANT

Company	**Angiomed U.S., Inc.**
Location	Anaheim, CA, US
Contact	RICHARD P MOHR
510(k) history	24 submissions · 24 cleared · 1986-1988