FDA 510(k) Premarket Notification

K834358 PARALLEL DUAL LUMEN SUBCLAVIAN CANNULA

CLEARED

Apr 2, 1984
111 days to decision

K834358 · Product code: **FIQ** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k834358/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, A-v Shunt (FIQ)
Date received	Dec 13, 1983
Decision date	Apr 2, 1984
Days to decision	111 days
Third-party review	No

APPLICANT

Company	**Vas-Cath of Canada , Ltd.**
Location	Mchenry, IL, US
510(k) history	19 submissions · 19 cleared · 1979-1988