FDA 510(k) Premarket Notification

K895506 FLEXXICON PLUS DUAL LUMEN DIALYSIS CATHETER

CLEARED

Feb 9, 1990
150 days to decision

K895506 · Product code: **LFJ** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k895506/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Subclavian (LFJ)
Date received	Sep 12, 1989
Decision date	Feb 9, 1990
Days to decision	150 days
Third-party review	No

APPLICANT

Company	**Vas-Cath, Inc.**
Location	Mississauga, Ontario, CA
Contact	H CREWE
510(k) history	27 submissions · 27 cleared · 1988-1998