FDA 510(k) Premarket Notification

K883768 VAS-CATH CATHETER REPAIR KIT

CLEARED

Nov 14, 1988
69 days to decision

K883768 · Product code: **LFJ** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k883768/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Subclavian (LFJ)
Date received	Sep 6, 1988
Decision date	Nov 14, 1988
Days to decision	69 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