FDA 510(k) Premarket Notification

K871487 VAS-CATH PERMANENT SINGLE LUMEN HEMODIALYSIS CATH

CLEARED

Jun 12, 1987
58 days to decision

K871487 · Product code: **LFJ** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k871487/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Subclavian (LFJ)
Date received	Apr 15, 1987
Decision date	Jun 12, 1987
Days to decision	58 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k871487/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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