

K834358 PARALLEL DUAL LUMEN SUBCLAVIAN CANNULAApr 2, 1984
111 days to decisionK834358 · Product code: **FIQ** · Gastroenterology & UrologySource: <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

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Device record: <https://www.510kdatabase.net/k834358/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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