

K812159 INTRAVASCULAR CATHETERAug 31, 1981
33 days to decisionK812159 · Product code: **FOZ** · General HospitalSource: <https://www.510kdatabase.net/k812159/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	Jul 29, 1981
Decision date	Aug 31, 1981
Days to decision	33 days
Third-party review	No

APPLICANT

Company	Pioneer Viggo, Inc.
Location	Mchenry, IL, US
510(k) history	6 submissions · 6 cleared · 1979-1982

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Device record: <https://www.510kdatabase.net/k812159/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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