FDA 510(k) Premarket Notification

K812159 INTRAVASCULAR CATHETER

CLEARED

Aug 31, 1981
33 days to decision

K812159 · Product code: **FOZ** · General Hospital
Source: 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