FDA 510(k) Premarket Notification

K910799 LANDMARK VENOUS ACCESS DEVICE

CLEARED

Mar 21, 1991
24 days to decision

K910799 · Product code: **FOZ** · General Hospital
Source: https://www.510kdatabase.net/k910799/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	Feb 25, 1991
Decision date	Mar 21, 1991
Days to decision	24 days
Third-party review	No

APPLICANT

Company	**Menlo Care, Inc.**
Location	Menlo Park, CA, US
Contact	JOCK M WALKER
510(k) history	31 submissions · 25 cleared · 1986-1997