FDA 510(k) Premarket Notification

K883980 LANDMARK VENOUS ACCESS DRIVE (ADDITIONAL SIZES)

CLEARED

Dec 5, 1988
75 days to decision

K883980 · Product code: **FOZ** · General Hospital
Source: https://www.510kdatabase.net/k883980/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	Sep 21, 1988
Decision date	Dec 5, 1988
Days to decision	75 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k883980/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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