FDA 510(k) Premarket Notification

K882100 VITACUFF(R) PERCUTANEOUS INFECT. CONTROL KIT

CLEARED

Jul 13, 1988
55 days to decision

K882100 · Product code: **LJS** · General Hospital
Source: https://www.510kdatabase.net/k882100/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Long-term Greater Than 30 Days (LJS)
Date received	May 19, 1988
Decision date	Jul 13, 1988
Days to decision	55 days
Third-party review	No

APPLICANT

Company	**Vitaphore Corp.**
Location	San Francisco, CA, US
Contact	SOPHIA PESOTCHINSKY
510(k) history	13 submissions · 12 cleared · 1986-1992