

**K882100 VITACUFF(R) PERCUTANEOUS INFECT. CONTROL
KIT**Jul 13, 1988
55 days to decisionK882100 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k882100/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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