

**K896046 VITAPATCH CATH. SECUREMENT DEVICE
W/CHLORHEXIDINE**Jan 11, 1990
86 days to decisionK896046 · Product code: **KMK** · General Hospital
Source: <https://www.510kdatabase.net/k896046/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Intravascular Catheter Securement (KMK)
Date received	Oct 17, 1989
Decision date	Jan 11, 1990
Days to decision	86 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/k896046/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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