

**K895993 VITAPATCH ANTIMICROBIAL CATHETER  
SECUREMENT DEVICE**Jan 11, 1990  
90 days to decisionK895993 · Product code: **KMK** · General Hospital  
Source: <https://www.510kdatabase.net/k895993/>**SUBMISSION DETAILS**

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|-----------------------|---|
| Decision              | Substantially Equivalent (Cleared)              |
| Submission type       | Traditional                                     |
| Device classification | Device, Intravascular Catheter Securement (KMK) |
| Date received         | Oct 13, 1989                                    |
| Decision date         | Jan 11, 1990                                    |
| Days to decision      | 90 days   |
| Third-party review    | No  |

**APPLICANT**

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

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k895993/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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