FDA 510(k) Premarket Notification

# K895920 VITAPATCH PIN PROTECTION DEVICE FOR PERCUTANEOUS

**CLEARED**

Jun 6, 1991
603 days to decision

K895920 · Product code: **KGX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k895920/

## SUBMISSION DETAILS

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|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Tape And Bandage, Adhesive (KGX) |
| Date received | Oct 11, 1989 |
| Decision date | Jun 6, 1991 |
| Days to decision | 603 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 |