

K895920 VITAPATCH PIN PROTECTION DEVICE FOR PERCUTANEOUSJun 6, 1991
603 days to decisionK895920 · Product code: **KGX** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k895920/>**SUBMISSION DETAILS**

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k895920/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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