FDA 510(k) Premarket Notification

K896046 VITAPATCH CATH. SECUREMENT DEVICE W/CHLORHEXIDINE

CLEARED

Jan 11, 1990
86 days to decision

K896046 · 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