FDA 510(k) Premarket Notification

K930225 K-LOK CATHETER SECUREMENT DEVICE

CLEARED

Apr 21, 1993
96 days to decision

K930225 · Product code: **KMK** · General Hospital
Source: https://www.510kdatabase.net/k930225/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Intravascular Catheter Securement (KMK)
Date received	Jan 15, 1993
Decision date	Apr 21, 1993
Days to decision	96 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**K-Lok, Inc.**
Location	Boca Raton, FL, US
Contact	GLENDA KALT
510(k) history	2 submissions · 2 cleared · 1993-1994