FDA 510(k) Premarket Notification

# K935252 K-LOK CATHETER SECUREMENT DEVICE MODIFICATION

**CLEARED**

Feb 18, 1994
134 days to decision

K935252 · Product code: **KMK** · General Hospital
Source: https://www.510kdatabase.net/k935252/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Device, Intravascular Catheter Securement (KMK) |
| Date received | Oct 7, 1993 |
| Decision date | Feb 18, 1994 |
| Days to decision | 134 days |
| Third-party review | No |
| Summary / Statement | Statement |

## APPLICANT

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