FDA 510(k) Premarket Notification

# K901969 ALERT BED ALARM SYSTEM

**CLEARED**

Sep 12, 1990
134 days to decision

K901969 · Product code: **KMI** · General Hospital
Source: https://www.510kdatabase.net/k901969/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Monitor, Bed Patient (KMI) |
| Date received | May 1, 1990 |
| Decision date | Sep 12, 1990 |
| Days to decision | 134 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Alertronics, Inc.** |
| Location | Tulsa, OK, US |
| Contact | RON WOOD |
| 510(k) history | 1 submissions · 1 cleared · 1990-1990 |