FDA 510(k) Premarket Notification

# K884106 SUPERVISORY TELEMETRY ARRHYTHMIA TERMINAL

**CLEARED**

Mar 6, 1989
158 days to decision

K884106 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k884106/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Detector And Alarm, Arrhythmia (DSI) |
| Date received | Sep 29, 1988 |
| Decision date | Mar 6, 1989 |
| Days to decision | 158 days |
| Third-party review | No |

## APPLICANT

| | |
|---|---|
| Company | **Siemens Medical Solutions USA, Inc.** |
| Location | Hoffman Estates, IL, US |
| Contact | HARRY K SCHWILL |
| 510(k) history | 778 submissions · 778 cleared · 1980-2026 |