

**K894460 SIEMENS SIRECUST 730, PATIENT MONITOR
W/ARRHYTHMIA**Oct 2, 1989
76 days to decisionK894460 · Product code: **DSI** · Cardiovascular
Source: <https://www.510kdatabase.net/k894460/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Jul 18, 1989
Decision date	Oct 2, 1989
Days to decision	76 days
Third-party review	No

APPLICANT

Company	Siemens Medical Electronics
Location	Danvers, MA, US
Contact	ROBERT R MURFITT
510(k) history	15 submissions · 15 cleared · 1988-1995

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k894460/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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