

**K862375 CARDIO PORT 500 24-H ECG AMBULATORY
MONITOR SYSTEM**Jul 1, 1986
7 days to decisionK862375 · Product code: **DSI** · Cardiovascular
Source: <https://www.510kdatabase.net/k862375/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Jun 24, 1986
Decision date	Jul 1, 1986
Days to decision	7 days
Third-party review	No

APPLICANT

Company	Grandcor Medical Systems
Location	Dayton, OH, US
Contact	SPERO M ALEX
510(k) history	8 submissions · 8 cleared · 1986-1991

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k862375/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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