FDA 510(k) Premarket Notification

K862375 CARDIO PORT 500 24-H ECG AMBULATORY MONITOR SYSTEM

CLEARED

Jul 1, 1986
7 days to decision

K862375 · 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