FDA 510(k) Premarket Notification

K860175 CD-200 CARDIAC MONITOR

CLEARED

Apr 15, 1986
88 days to decision

K860175 · Product code: **DPS** · Cardiovascular
Source: https://www.510kdatabase.net/k860175/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Jan 17, 1986
Decision date	Apr 15, 1986
Days to decision	88 days
Third-party review	No

APPLICANT

Company	**Cardio Display Corp.**
Location	Westbury, NY, US
Contact	MILAN KESLER
510(k) history	2 submissions · 2 cleared · 1986-1986