FDA 510(k) Premarket Notification

K863159 Q4000 AND Q3040 ELECTROCARDIOGRAPH MONITOR

CLEARED

Oct 6, 1986
49 days to decision

K863159 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k863159/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Aug 18, 1986
Decision date	Oct 6, 1986
Days to decision	49 days
Third-party review	No

APPLICANT

Company	**Quinton, Inc.**
Location	Mchenry, IL, US
Contact	RON R DUCK
510(k) history	164 submissions · 160 cleared · 1976-2003