

**K863159 Q4000 AND Q3040 ELECTROCARDIOGRAPH MONITOR**Oct 6, 1986  
49 days to decisionK863159 · 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	<b>Quinton, Inc.</b>
Location	Mchenry, IL, US
Contact	RON R DUCK
510(k) history	164 submissions · 160 cleared · 1976-2003

---

**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k863159/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 20, 2026