

**K882978 Q4000 OPTION FOR Q4000**Nov 1, 1988  
109 days to decisionK882978 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k882978/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Jul 15, 1988
Decision date	Nov 1, 1988
Days to decision	109 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/k882978/>; 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