FDA 510(k) Premarket Notification

# K882978 Q4000 OPTION FOR Q4000

**CLEARED**

Nov 1, 1988
109 days to decision

K882978 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k882978/

## SUBMISSION DETAILS

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|---|---|
| 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

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