

K820438 QUINTON Q2000 ETT MONITORING SYSTEMApr 29, 1982
71 days to decisionK820438 · Product code: **DSI** · Cardiovascular
Source: <https://www.510kdatabase.net/k820438/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Feb 17, 1982
Decision date	Apr 29, 1982
Days to decision	71 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k820438/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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